

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
2 December 2004 (02.12.2004)

PCT

(10) International Publication Number
WO 2004/104681 A1

(51) International Patent Classification⁷: **G02F 1/13363**,
G08G 64/04

(21) International Application Number:
PCT/US2004/013676

(22) International Filing Date: 3 May 2004 (03.05.2004)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
10/439,541 16 May 2003 (16.05.2003) US

(71) Applicant (for all designated States except US): **EASTMAN KODAK COMPANY** [US/US]; 343 State Street, Rochester, New York 14650-2201 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **ELMAN, James Frank** [US/US]; 2543 Huber Road, Fairport, New York 14450 (US). **ISHIKAWA, Tomohiro** [JP/US]; 252 Mulberry Street, Rochester, New York 14620 (US). **MASSA, Dennis Jon** [US/US]; 35 Cambric Circle, Pittsford, New York 14534 (US). **YACOBUCCI, Paul Daniel** [US/US]; 5 Windsor Park, Rochester, New York 14624 (US). **FALKNER, Catherine Ann** [US/US]; 236 Wayfaring Lane, Rochester, New York 14612 (US).

(74) Common Representative: **EASTMAN KODAK COMPANY**; 343 State Street, Rochester, New York 14650-2201 (US).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

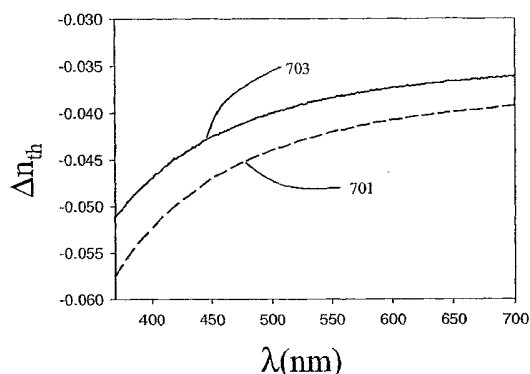
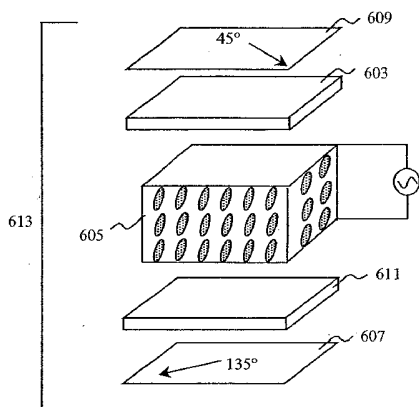
(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Declarations under Rule 4.17:

— as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii)) for the following designations AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM,

[Continued on next page]

(54) Title: COMPENSATION FILMS FOR LCDS



(57) Abstract: Disclosed is an optical compensation film (603, 611) for Liquid Crystal Displays comprising an amorphous polymer layer disposed on a substrate wherein said amorphous polymer layer has an out-of-plane birefringence $\Delta n_{th}(\lambda)$ (701, 703) at a wavelength λ which is a monotonic increasing function (i.e. it has a proper dispersion) that satisfies both conditions (1) and (2): Condition (1) $\Delta n_{th}(\lambda)$ is more negative than -0.005 throughout the range of $370\text{nm} < \lambda < 700\text{nm}$; Condition (2) $\Delta n_{th}(\lambda_1)$ is more negative than $\Delta n_{th}(\lambda_2)$, for $\lambda_1 < \lambda_2$ throughout the range of $370\text{nm} < \lambda < 700\text{nm}$, and wherein the polymer does not have chromophores off the backbone.



- ZW, ARIPO patent (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG)
- as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii)) for all designations

Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

COMPENSATION FILMS FOR LCDS

FIELD OF THE INVENTION

The present invention relates to an optically transparent
 5 compensation film for liquid crystal displays (LCD) comprising an amorphous
 polymer layer with negative out-of-plane birefringence exhibiting proper
 dispersion. The invention also provides a process for making such a
 compensation film.

10 BACKGROUND OF THE INVENTION

The following terms have the definitions as stated below.

Optic axis herein refers to the direction in which propagating light does not see
 birefringence.

A-plate and C-plate herein are the plates in which the optic axis is in the plane of
 15 the plate and perpendicular to the plate.

Polarizer herein refers to elements that polarize an electromagnetic wave.

In-plane phase retardation, R_{in} , of a film **201** shown in FIG. 1 is a quantity defined
 by $(n_x - n_y)d$, where n_x and n_y are indices of refraction in the direction of x and y ,
 respectively and x - y plane is parallel to the plane **203** of the film. d is a thickness
 20 of the film in z -direction. The quantity $(n_x - n_y)$ is referred as in-plane
 birefringence. Both of these in-plane quantities will be treated as absolute values
 with no preferred sign convention.

Out of-plane phase retardation, R_{th} , of a film **201** shown in FIG. 1, herein, is a
 quantity defined by $[n_z - (n_x + n_y)/2]d$. n_z is the index of refraction in z -direction.

25 The quantity $[n_z - (n_x + n_y)/2]$ is referred as out-of-plane birefringence, Δn_{th} . If
 $n_z > (n_x + n_y)/2$, Δn_{th} is positive, thus the corresponding R_{th} is also positive. If
 $n_z < (n_x + n_y)/2$, Δn_{th} is negative and R_{th} is also negative.

Amorphous herein means a lack of long-range order. Thus an amorphous polymer
 does not show long-range order as measured by techniques such as X-ray
 30 diffraction.

Liquid crystals are widely used for electronic displays. In these
 display systems, a liquid crystal cell is typically situated between a pair of

polarizers. An incident light polarized by the polarizer passes through a liquid crystal cell and is affected by the molecular orientation of the liquid crystal, which can be altered by the application of a voltage across the cell. The altered light goes into the second polarizer. By employing this principle, the transmission of light
5 from an external source, including ambient light, can be controlled. The energy required to achieve this control is generally much less than required for the luminescent materials used in other display types such as cathode ray tubes (CRT). Accordingly, LCD technology is used for a number of electronic imaging devices, including but not limited to digital watches, calculators, portable
10 computers, electronic games, and televisions for which light-weight, low-power consumption and long-operating life are important features.

Contrast, color reproduction, and stable gray scale intensities are desirable attributes for electronic displays, which employ LCD technology. The primary factor limiting the contrast of a LCD is the propensity of the light to
15 "leak" through liquid crystal elements or cells, which are in the dark or "black" pixel state. Furthermore, this leakage and hence the contrast of a liquid crystal display are also dependent on the direction from which the display is viewed. Typically the optimum contrast is observed only within a narrow viewing angle range centered about the normal incidence to the display and falls off rapidly as
20 viewing direction moves away from the display normal. In color LCDs, this leakage problem not only degrades the contrast but also causes color or hue shifts with an associated degradation of color reproduction. There are various modes of LCDs. Twisted Nematic (TN) LCDs are liquid crystal displays in which optic axis of liquid crystal rotates 90° in the azimuthal angle across the liquid crystal cell
25 thickness direction when no field is applied. With a sufficiently large applied field, the liquid crystal optic axis becomes perpendicular to the liquid crystal cell plane except in the vicinity of cell bounding plate. In the vicinity of the cell bounding plate, the liquid crystal optic axis deviates from the cell normal direction. Vertically Aligned (VA) LCDs have liquid crystal optic axis that is
30 substantially perpendicular to the liquid crystal cell plane without an applied field. This state corresponds to a dark state of the displays. With applied field, liquid crystal optic axis tilt away from the cell normal direction. Optically Compensated

Bend (OCB) LCDs are liquid crystal displays based on the symmetric bow-shape bend state of liquid crystal optic axis. The bow-shape bend state occurs in the plane perpendicular to the liquid crystal cell plane and the state of bend is symmetric around the mid point in the cell thickness direction. In-plane switching (IPS) LCDs are liquid crystal display in which the field to change the direction of the liquid crystal optic axis is applied in the plane of the liquid crystal cell. In IPS LCDs, the liquid crystal optic axis changes its direction while remaining substantially in the plane of the liquid crystal cell.

LCDs are quickly replacing CRT's as monitors for desktop computers and other office or household appliances. It is also expected that the number of LCD television monitors with a larger screen size will sharply increase in the near future. However, unless problems of viewing angle dependence such as coloration, degradation in contrast, and an inversion of brightness are solved, the replacement of the traditional CRT by LCDs will be limited.

One of the common methods to improve LCD viewing angle characteristic is to use compensation films. Situated between a polarizer and a liquid crystal cell, a compensation film annuls the phase retardation imposed on the propagating light by the liquid crystal cell. Several LCD modes, with or without an applied field, exhibit positive C-plate symmetry that can be compensated by a compensation film with negative C-plate property. As is well known to those who are skilled in the art, TN and OCB Liquid Crystal Cell show C-plate symmetry in the center portion of the cell with sufficiently high, applied electric field. The VA Liquid Crystal Cell exhibits C-plate symmetry in the state without an applied field. This approximate positive C-plate state of a liquid crystal cell gives a dark state, if it is placed between the crossed polarizers. Here, crossed polarizers mean that transmission (or absorption) axes of two polarizers form an angle of $90\pm 10^\circ$. The ray propagating perpendicular to the liquid crystal cell essentially does not see birefringence. That is the reason why in the normal viewing direction one has the highest contrast ratio in some modes of LCDs. On the other hand, obliquely propagating light rays see the phase retardation and this leads to light leakage in the dark state and degrades the contrast ratio. FIG. 2 schematically shows the principle of compensation. Ellipsoid 301 represents the

positive C-plate that approximates liquid crystal cell **305** having $n_{xlc} = n_{yic} = n_{olic}$, $n_{zlc} = n_{elic}$ where $n_{elic} > n_{olic}$. n_{elic} and n_{olic} are extraordinary and ordinary indices of liquid crystal. Compensation film **307** with negative R_{th} is shown by the ellipsoid **309**. Here we have $(n_x + n_y)/2 > n_z$, thus giving a negative R_{th} . For the rays **311**

5 obliquely propagating with an incident angle ϕ through the liquid crystal cell **305**, the positive phase retardation from liquid crystal cell **305** is canceled by the negative phase retardation of the compensation film **307**. Therefore, one can effectively prevent the light leakage caused by the birefringence of the liquid crystal cell in the oblique direction. Thus, compensation film for various modes of

10 LCDs must have a negative R_{th} and simple methods to obtain this negative R_{th} are highly desirable for optical compensation of LCDs.

One of the essential attributes of LCD compensation is the wavelength dependence (or dispersion) of the phase retardation and birefringence versus the wavelength of light (λ). It is important to achieve a proper dark state

15 without color shift. Phase retardation is directly proportional to the birefringence, and their dispersion shape (proper or reverse) is likewise related. "Proper" dispersion is such that the absolute value of phase retardation and birefringence increases toward shorter λ (i.e. towards the ultraviolet region). Conversely, materials with "reverse" dispersion have smaller absolute value of phase

20 retardation and birefringence in the shorter λ region. Typical liquid crystal cells use materials with proper dispersion, which show larger positive values of birefringence and phase retardation toward shorter λ . For the negative birefringence case (the compensation film), proper dispersion gives larger

25 negative value of birefringence and phase retardation at shorter λ . Conversely, negative birefringence with reverse dispersion (i.e. less desirable compensation films) exhibits less negative birefringence and phase retardation for light with shorter λ . The phase retardation dispersion of the liquid crystal cell and the compensation film have to be the same kind to properly cancel the phase

30 retardation of a liquid crystal cell by compensation film. That is, if the liquid crystal phase retardation assumes larger positive value at shorter λ (i.e. positive birefringence with proper dispersion), the compensation films have to have larger

negative value (i.e. negative birefringence with proper dispersion) of R_{th} (or Δn_{th}) for good compensation.

FIG. 3 explains the concept. The liquid crystal cell in the positive C-plate state has a positive out-of-plane birefringence $\Delta n_{lc} = (n_{e_{lc}} - n_{o_{lc}})$. The phase retardation to be compensated by a compensation film with negative out-of-plane birefringence and phase retardation is $R_{th-lc} = d_{lc} \Delta n_{lc}$ (where d_{lc} is a thickness of a liquid crystal cell). The curves show the wavelength dependence of $R_{th-lc} = d_{lc} \Delta n_{lc}$, **401** and $R_{th} = d \Delta n_{th} = d[nz - (nx + ny)/2]$ of the compensation films, **403**, **407**. A typical liquid crystal is positively birefringent and has proper birefringence dispersion. Namely, the birefringence Δn_{lc} increases toward the shorter wavelength λ . Therefore, the phase retardation R_{th-lc} assumes a larger positive value at the shorter λ as shown by the curve **401**. To counterbalance it, the compensation film is required to have also proper phase retardation dispersion shown by the curve **403**. Films with the property shown by the curve **403** have a larger negative value of Δn_{th} and R_{th} at the shorter λ . On the other hand, curve **407** shows the compensation film of reverse dispersion. In this case, the birefringence of the film Δn_{th} and R_{th} has a smaller negative value at the shorter λ . Thus, proper cancellation of the phase retardation of a liquid crystal cell (a positive C-plate) cannot be achieved by a film with reverse dispersion for a wide range of wavelengths. Another important aspect of the compensation films is the transparency. At each wavelength λ , percent transmission of the film $T(\lambda)$ can be measured. Percent transmission $T(\lambda)$ gives the intensity percentile of the transmitted light with respect to the incident light in the film normal direction. The average transmission in the wavelength range $\lambda_1 \leq \lambda \leq \lambda_2$ is defined as $T_{av} =$

$$\frac{\int_{\lambda_1}^{\lambda_2} T(\lambda) d\lambda}{\lambda_2 - \lambda_1}.$$

Optically transparent film is a film such that $T_{av} \geq 90\%$ for $\lambda_1 = 400\text{nm}$

and $\lambda_2 = 700\text{nm}$. Such a film is preferred, as it does not compromise the brightness of the image on LCDs. Several means of obtaining negative R_{th} have been suggested. Sergan et al. discusses the crossed A-plate as a replacement of the negative C-plate ("Two Crossed A-plates as an Alternative to a Negative C-plate",

Society of Information Display 2000, pp. 838-841). Two A-plates are placed on top of each other with their optic axes forming 90° . They showed that crossed A-plates function approximately as a negative-C plate and successfully compensated the VA Liquid Crystal Cell. This method, however, involves cumbersome process of laminating two films with A-plate property to form one compensation film of negative-C behavior. Further, it is known to those who skilled in the art that crossed A-plates do not show negative C-plate behavior to rays with a small incident angle ϕ . Thus it is not a desirable method of obtaining negative R_{th} .

JP 1999-95208 discloses the use of a swellable inorganic clay layer in a crosslinked organic matrix to generate negative R_{th} . The disclosed method enables continuous means of manufacturing films with negative C-plate character. However, the resulting film gives the reverse dispersion, represented by a curve 407. Thus it is not suitable as a compensation film.

Li et al. ("Polyimide film as negative birefringent compensators for normally white twisted nematic liquid crystal displays", Polymer, Volume 37, pp 5321-5325, (1996)) disclosed a polyimide layer formed by spin-casting or dip-emersion method. The film shows proper dispersion. This paper describes several polyimides that could be used as compensation films with negative R_{th} . However, the T_{av} is less than 90%. Also, as is well known to those who skilled in the art, polyimides generally suffer from a yellow-orange coloration. Thus, they are not desirable to be used as a compensation film because the color of the film would shift the hue of the images on the LCD's.

The use of a biaxially stretched cellulose ester film as a compensation film is disclosed in JP 2002-236216. Biaxially stretched cellulose ester film with large negative R_{th} and positive R_{in} is used as a substrate on which optically anisotropic layer with O-plate character is disposed. This biaxially stretched cellulose ester film (according to the disclosure) exhibits sufficient negative R_{th} to be useful as a compensation film. However, it possesses a reverse dispersion in Δn_{th} , and thus in R_{th} .

Prior arts offer methods of obtaining films with sufficiently large negative R_{th} value. However, prior art films do not offer films with proper dispersion in negative R_{th} and Δn_{th} that can be easily manufactured and have a

high T_{av} ($T_{av} \geq 90\%$, that is optically transparent). Also, it is highly desirable to obtain thinner compensation film with large negative R_{th} value. This enables overall LCD packages, comprising liquid crystal cell, compensation films, and polarizers to be slimmer. Thus, a desired method would offer a compensation film with sufficiently large negative R_{th} and the desired, proper retardation/birefringence dispersion, without significant increase in the display thickness.

SUMMARY OF THE INVENTION

The invention provides an optical compensation film for Liquid Crystal Displays that has a negative R_{th} with a proper dispersion. It comprises an amorphous polymer layer disposed on a substrate. The out-of-plane birefringence $\Delta n_{th}(\lambda)$ of the amorphous polymer at a wavelength λ satisfies conditions (1) and (2), respectively:

Condition (1)

$\Delta n_{th}(\lambda)$ is more negative than -0.005 throughout the range or $370\text{nm} < \lambda < 700\text{nm}$;

Condition (2)

$\Delta n_{th}(\lambda_1)$ is more negative than $\Delta n_{th}(\lambda_2)$, for $\lambda_1 < \lambda_2$ throughout the range of $370\text{nm} < \lambda < 700\text{nm}$.

Thus, the compensation film comprises a layer with large, negative values of Δn_{th} . This amorphous polymer layer enables an increase in the negative value of R_{th} of the compensation film, without a significant thickening of the film. The invention also includes a Liquid Crystal Display including the compensation film and a process for forming the compensation film.

BRIEF DESCRIPTION OF THE DRAWINGS

While the specification concludes with claims particularly pointing out and distinctly claiming the subject matter of the present invention, it is believed that the invention will be better understood from the following description when taken in conjunction with the accompanying drawings, wherein:

FIG.1 shows the film with thickness d and x-y-z coordinate system attached to the film;

FIG. 2 shows the principle of compensation of a liquid crystal cell with positive C-plate character with a compensation film with negative R_{th} ;

5 FIG. 3 shows a curve of proper dispersion of liquid crystals and curves for compensation films with proper and reverse dispersion;

FIG. 4A, FIG. 4B and FIG. 4C are elevation schematics of the examples of compensation film;

10 FIG. 5A and FIG. 5B show LCD's with one and two compensation film, respectively;

FIG. 6 shows proper dispersion in birefringence curves of an amorphous polymer layer;

FIG. 7 shows the proper dispersion in phase retardation of an exemplary compensation film according to the invention;

15 FIG. 8 shows the reverse dispersion in phase retardation curve of comparative example;

DETAILED DESCRIPTION OF THE INVENTION

20 Reference will now be made to the drawings in which the various elements of the present invention will be given numerical designations and in which the invention will be discussed so as to enable one skilled in the art to make and use the invention. It is to be understood that elements not specifically shown or described may take various forms well known to those skilled in the art.

25 The invention provides an optical compensation film for Liquid Crystal Displays that has a negative R_{th} with a proper dispersion. It comprises an amorphous polymer layer disposed on the substrate. The out-of-plane birefringence $\Delta n_{th}(\lambda)$ of the amorphous polymer at a wavelength λ that satisfies conditions (1) and (2), respectively:

Condition (1)

30 $\Delta n_{th}(\lambda)$ is more negative than -0.005 throughout the range of $370\text{nm} < \lambda < 700\text{nm}$;

Condition (2)

$\Delta n_{th}(\lambda_1)$ is more negative than $\Delta n_{th}(\lambda_2)$, for $\lambda_1 < \lambda_2$ throughout the range of
370nm < λ < 700nm.

5

FIGS. 4A, 4B and 4C show the elevation views of exemplary structures of the compensation films. In compensation film **501**, an amorphous polymer layer **503** is disposed on the substrate **505**. The polymer layer **503** satisfies both conditions (1) and (2). FIG. 4B shows the compensation film **511** where an additional layer **507** is disposed on the polymer layer **503**. The layer **507** can be, for example, an antiglare coating or anti-reflection coating. It can also be a hard-coat layer that mechanically protects the amorphous polymer layer **503**. The order can change such that the layer **507** is disposed on the substrate followed by **503** such as the one **513** shown in FIG. 4C. In this case, **507** may have a function of buffer layer between the substrate **505** and the amorphous polymer layer **503**. The buffer layer can promote the adhesion of the layer **503** to the substrate **505**. In some other cases, it prevents the diffusion of chemical contamination from the substrate **505** to the polymer layer **503**, thus acting as a barrier layer. Also, it may be a compliance layer to prevent the cracking of the polymer layer **503**.

The substrate **505** can be optically isotropic or anisotropic. As is well known in the art, optical materials may have up to three different principal indices of refraction and can be classified as either isotropic or anisotropic based on the relationships of these indices. When all three of its principal indices are equal, a material is said to be isotropic. When anisotropic, a material can be either uniaxial, or biaxial. If two principal indices are equal, a material is called uniaxial. A uniaxial material is uniquely characterized, as having an ordinary index, referred as n_o , an extraordinary index n_e and two angles describing the orientation of its optical axis, the axis of n_e . When n_e is greater than n_o , a uniaxial material is positively birefringent. When n_e is smaller than n_o , a uniaxial material is negatively birefringent. Controlling birefringent behavior is particularly useful in the fabrication and application of optical films. When all

three refractive indices are different, a material is said to be biaxial, uniquely specified by its principal indices n_x , n_y , n_z , and three orientation angles. Some of biaxial materials show weak biaxiality meaning that two of their three principal indices are very close, which is often considered equally as the ordinary refractive index for a uniaxial material.

For an ease of the manufacturing, it is preferable to have a flexible substrate, particularly polymeric substrate. Typical polymeric substrate is triacetylcellulose (TAC). The optical property of TAC is close to the uniaxial and can be approximated as a negative C-plate, thus TAC has negligible R_{in} and negative R_{th} . In other cases, biaxially stretched TAC is used for a substrate. In this case, the substrate has finite R_{in} and R_{th} . TAC has reverse dispersion in out-of plane birefringence Δn_{th} . This reverse dispersion of TAC, however, is compensated by proper dispersion in Δn_{th} of the amorphous polymer layer **503**. Low birefringent polymeric material such as cyclic polyolefin can also be a substrate.

The amorphous polymer layer **503** is coated from a solution containing a polymer that yields large negative Δn_{th} upon solvent coating. The large negative value of Δn_{th} with proper dispersion will dominate the total negative R_{th} of the compensation film and its kind of dispersion. Thus, by disposing a polymer layer **503** on top of a substrate (with negative R_{th} and reverse dispersion), the overall negative R_{th} will increase and the dispersion shape will become "proper". To produce an amorphous polymer layer with negative Δn_{th} satisfying the condition (1) and (2), polymers that contain non-visible chromophore groups such as vinyl, carbonyl, amide, ester, carbonate, sulfone, azo, and aromatic groups (i.e. benzene, naphthalate, biphenyl, bisphenol A) in the polymer backbone will be used, such as polyesters, polycarbonates, polyetherimides, and polythiophenes. Preferably, polymers to be used in the layer **503** do not have chromophores off of the backbone. An example of such an undesirable polymer with chromophores in and off the backbone would be polyarylates possessing the fluorene group. The glass transition temperature (T_g) of the polymers used in the amorphous layer is a significant factor. It should be

above 180° C to achieve the desired results. The thickness of the amorphous polymer layer **503** is determined by the necessary value of R_{th} that is at least more negative than -20nm. Typically it should be from -600nm to -60nm.

Conveniently it should be from -500nm to -100nm. Desirably it should be from -400nm to -150nm. The thickness of the polymer layer **503** should be less than 30 μ m. Typically it should be from 0.1 μ m to 20 μ m. Conveniently it should be from 1.0 μ m to 10 μ m. Compensation films with structures other than shown in FIG. 4A, FIG. 4B and FIG. 4C are possible.

The overall thickness of the compensation film is important to keep overall LCD package thickness within the reasonable range. The combined thickness of the compensation film should be less than 115 μ m. Typically it should be from 40 μ m to 105 μ m. Desirably it should be from 40 μ m to 100 μ m. The compensation film is optically transparent, namely, $T_{av} \geq 90\%$.

FIG. 5A shows a schematic liquid crystal display **601** where **603** is a single compensation film is placed on one side of the liquid crystal cell **605**. **607** is a polarizer, and **609** is a second polarizer. The transmission axes for the polarizers **607** and **609** form $90^\circ \pm 10^\circ$ angle relative to each other. The angles of their transmission axes are denoted as 45° and 135° . However, other angles are possible depending on the kind of liquid crystal display **601** and this is obvious to those who skilled in the art. Note that liquid crystal cell **605** is the electrically switchable liquid crystal cell with the liquid crystals confined between two glass plates.

FIG. 5B shows another schematic liquid crystal display **613** where there are two compensation films **603**, **611** placed on both sides of the liquid crystal cell **605**. **607** is a polarizer and **609** is a second polarizer. The transmission axes for the polarizers **607** and **609** form $90^\circ \pm 10^\circ$ angle relative to each other. The angles of their transmission axes are denoted as 45° and 135° . However, other angles are possible depending on the kind of liquid crystal display **601** and this is obvious to those who skilled in the art. Note that **605** is the electrically switchable liquid crystal cell with the liquid crystals confined between two glass plates.

Compared to the prior art, embodiments of the present invention provide the compensation film with negative R_{th} with proper dispersion that is optically transparent. Also, it provides a method of generating films with sufficiently large negative value of R_{th} with a thickness that is much smaller than the conventional films such as TAC. This is possible because of the large negative value of Δn_{th} of the amorphous polymer layer. Thus the invention enables enhanced optical compensation in a relatively thin ($<115\mu m$) structure that can be easily manufactured.

The present invention is further illustrated by the following non-limiting examples of its practice.

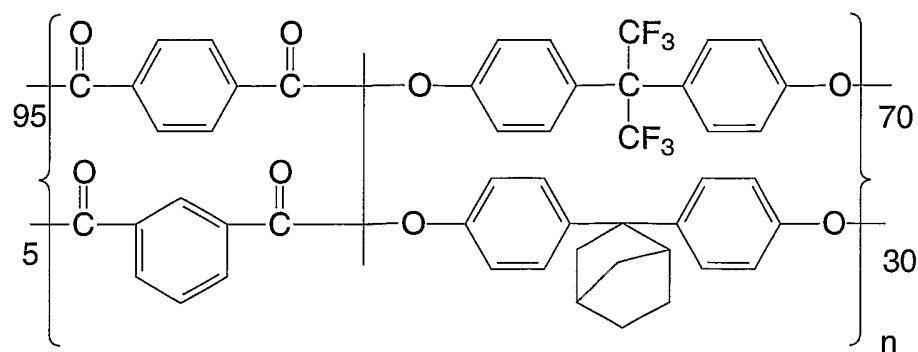
Examples:

The aromatic polyesters used herein can be prepared using any suitable or conventional procedure. The procedure used herein followed that outlined by P. W. Morgan in Condensation Polymers: By Interfacial and Solution Methods, Interscience, New York City, N.Y. (1965).

Example A:

Polymer A (synthesis):

To a stirred mixture of 4,4'-hexafluoroisopropylidenediphenol (23.53 g, 0.07 mole), 4,4'-(2-norbornylidene) bisphenol (8.4 g, 0.03 mole) and triethylamine (22.3 g, 0.22 mole) in methyl ethyl ketone (100 mL) at $10^{\circ}C$. was added a solution of terephthaloyl chloride (19.29 g, 0.095 mole) and isophthaloyl chloride (1.02 g, 0.005 mole) in methyl ethyl ketone (60 mL). After the addition, the temperature was allowed to rise to room temperature and the solution was stirred under nitrogen for 4 hours, during which time triethylamine hydrochloride precipitated in a gelatinous form and the solution became viscous. The solution was then diluted with toluene (160 mL) and washed with dilute hydrochloric acid, (200 mL of 2% acid) followed three times by water (200 mL). The solution was then poured into ethanol with vigorous stirring, and a white bead like polymer precipitated, collected and dried at $50^{\circ}C$. under vacuum for 24 hours. The glass transition temperature of this polymer was measured by differential scanning calorimetry to be $265^{\circ}C$.



Poly(4,4'-hexafluoroisopropylidene-bisphenol-co- 4,4'-(2-norbornylidene) bisphenol) terephthalate-co-isophthalate.

5

Polymer A

Polymer A was spun cast (8% solids in 80% propylacetate 20% toluene) onto a glass slide, and was analyzed with an ellipsometer (model M2000V, J.A. Woollam Co.) at 550nm wavelength to obtain the R_{th} and the thickness of the layer of polymer A. These values are listed in TABLE Ia.

TABLE Ia

Polymer A Layer thickness (μm)	R_{th} , Out-of-Plane Retardation (nm)
1.07	-45

R_{th} was similarly measured at other wavelengths and thus the Δn_{th} was calculated by knowing retardation R_{th} and layer thickness as shown in TABLE Ib..

15

TABLE Ib

Wavelength (nm)	out-of-plane birefringence Δn_{th} (unit dimension)
370	-0.057
550	-0.042
700	-0.039

The layer of polymer A also did not show any sign of a long-range order therefore the layer was determined to be comprised of an amorphous

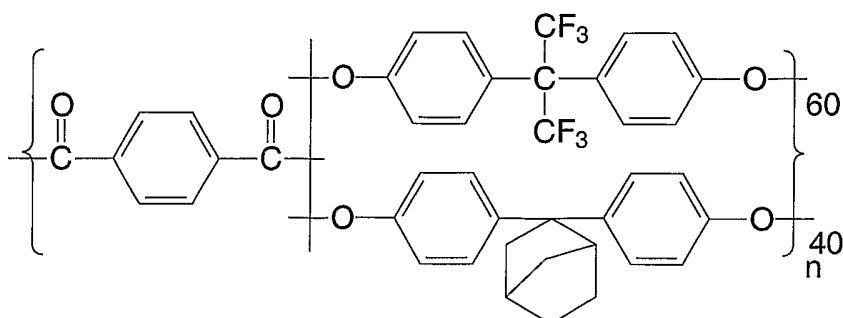
polymer. The dispersion curve **701** corresponding to Δn_{th} of a polymer A layer is shown in FIG. 7. It shows larger negative value of Δn_{th} at shorter wavelength λ . Thus, an amorphous polymer layer prepared according to the above mentioned method exhibits proper dispersion. Δn_{th} is more negative than -0.005 at the
5 wavelengths between 370 and 700nm.

Additionally, a layer of polymer A is disposed on a triacetylcellulose (TAC) film (thickness $80\mu m$). The polymer A layer had a thickness of $3.6\mu m$ as measured by ellipsometry. FIG. 7 shows the dispersion curve **801** of R_{th} of the resulting compensation film. The compensation film shows
10 proper dispersion. Transmission was measured for $400nm \leq \lambda \leq 700nm$ and $T_{av} \geq 90\%$ was obtained.

Example B:

Polymer B (synthesis):

15 To a stirred mixture of 4,4'-hexafluoroisopropylidenediphenol (23.53 g, 0.07 mole), 4,4'-(2-norbornylidene) bisphenol (8.4 g, 0.03 mole) and triethylamine (22.3 g, 0.22 mole) in dichloromethane (100 mL) at $10^\circ C$. was added a solution of terephthaloyl chloride (19.29 g, 0.095 mole) in dichloromethane (60 mL). After the addition, the temperature was allowed to rise
20 to room temperature and the solution was stirred under nitrogen for 4 hours, during which time triethylamine hydrochloride precipitated in a gelatinous form and the solution became viscous. The solution was then diluted with toluene (160 mL) and washed with dilute hydrochloric acid, (200 mL of 2% acid) followed three times by water (200 mL). The solution was then poured into ethanol with
25 vigorous stirring, and a white bead like polymer precipitated, collected and dried at $50^\circ C$. under vacuum for 24 hours. The glass transition temperature of this polymer was measured by differential scanning calorimetry to be $274^\circ C$



Poly(4,4'-hexafluoroisopropylidene-bisphenol-co- 4,4'-(2-norbornylidene)
bisphenol) terephthalate.

5

Polymer B

Polymer B was spun cast onto a glass slide (10% solids in 50% methyl ethyl ketone 50% toluene), and then was removed from this substrate. R_{th} and the polymer layer thickness of the layer thus made were measured with an
10 ellipsometer (model M2000V, J.A. Woollam Co.) at 550nm wavelength. Results are shown in the TABLE IIa

TABLE IIa

Polymer B Layer thickness (μm)	R_{th} , Out-of-Plane Retardation (nm)
4.02	-154

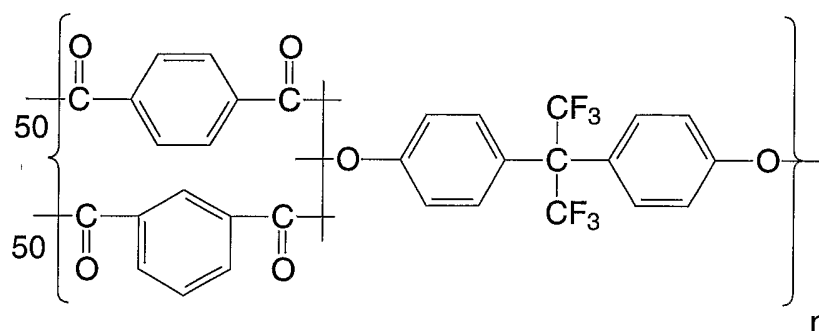
Retardation was similarly measured at other wavelengths (show in TABLE IIb)
15 and thus the Δn_{th} was calculated by knowing retardation and layer thickness.

Table IIb

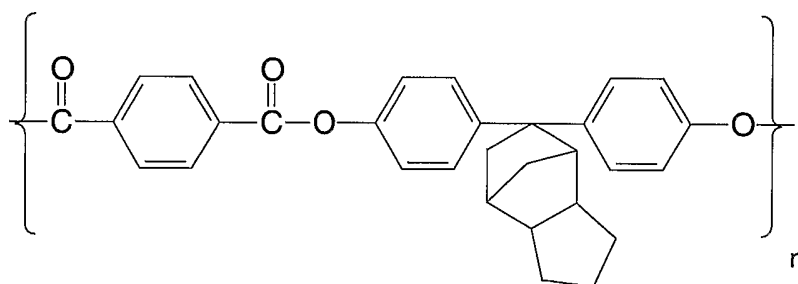
Wavelength (nm)	out-of-plane birefringence Δn_{th} (unit dimension)
370	-0.051
550	-0.038
700	-0.036

The layer of polymer B did not show any sign of a long-range order. Therefore the layer was determined to be comprised of an amorphous polymer. The dispersion curve **703** of Δn_{th} of polymer B is shown in FIG. 7. It shows larger negative value of Δn_{th} at shorter wavelength λ . Thus, an amorphous polymer layer prepared according to the above mentioned method exhibits proper dispersion. Δn_{th} is more negative than -0.005 at the wavelengths between 370 and 700nm.

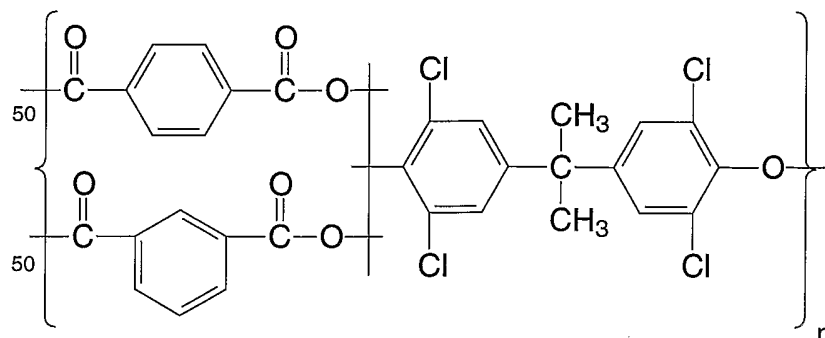
Other specific polymers that could be used include:



Poly(4,4'-hexafluoroisopropylidene-bisphenol) terephthalate-co-isophthalate
Polymer C



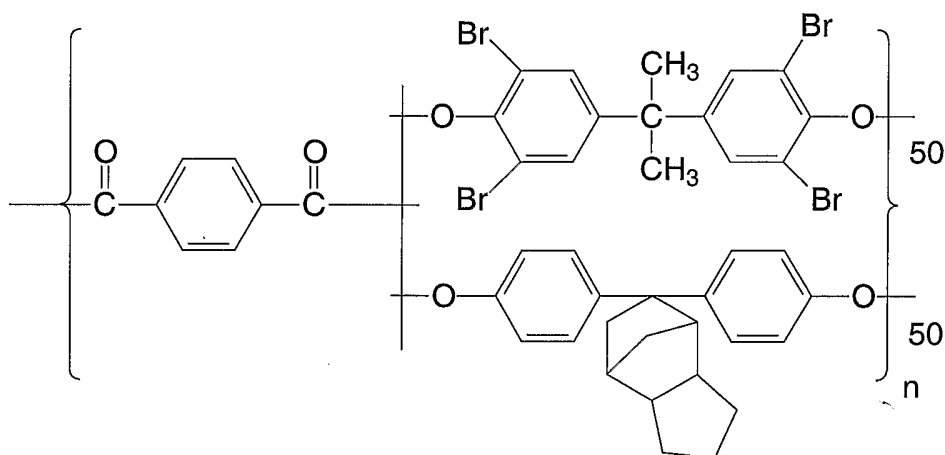
Poly(4,4'-hexahydro-4,7-methanoindan-5-ylidene bisphenol) terephthalate.
Polymer D



Poly(4,4'-isopropylidene-2,2',6,6'-tetrachlorobisphenol) terephthalate-co-isophthalate

5

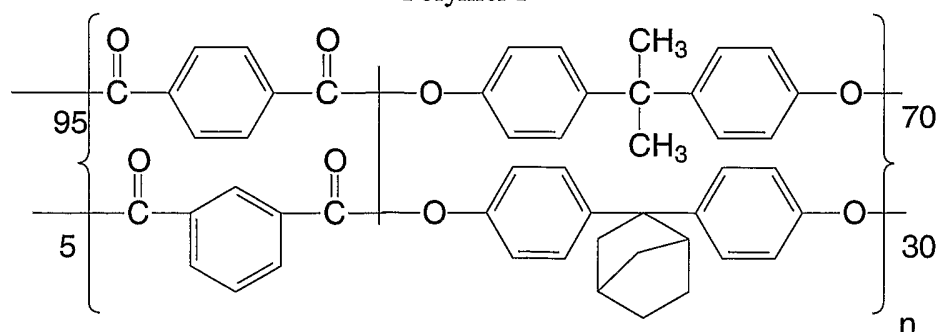
Polymer E



10

Poly(4,4'-hexahydro-4,7-methanoindan-5-ylidene)-bisphenol-co-(4,4'-isopropylidene-2,2',6,6'-tetrabromo)-bisphenol terephthalate.

Polymer F



15

Poly(4,4'-isopropylidene-bisphenol-co-4,4'-(2-norbornylidene) bisphenol) terephthalate-co-isophthalate

Polymer G

A series of polymers were analyzed for their glass transition temperatures and out of plane birefringence values Δn_{th} . It was found that the more desirable polymers for this invention had glass transition temperatures above 180°C. Those with lower glass transition temperatures were found to, generally, have out-of-plane birefringence Δn_{th} values less negative than -0.005 .

Comparative Example:

R_{th} of a cellulose ester film (thickness: $110\mu m$ as measured by a micrometer) was measured with an ellipsometer (model M2000V, J.A. Woollam Co.) at wavelengths between 400 to 700nm. Values of R_{th} at three wavelengths are listed in the Table III.

Table III

Wavelength (nm)	Out-of-plane retardation R_{th} (nm)
400	-244
550	-257
700	-266

15

The dispersion curve **901** of R_{th} of the comparative example film is shown in FIG. 8. It shows smaller negative value of R_{th} at shorter wavelength λ , thus the comparative example film exhibits reverse dispersion. From the measured thickness and R_{th} , Δn_{th} was calculated. Δn_{th} is less negative than -0.005 at the wavelength between 370 and 700nm. Note that this film does not show enough negative birefringence and does not have the desired dispersion sign.

The invention has been described in detail with particular reference to certain preferred embodiments thereof, but it will be understood that variations and modifications can be effected within the scope of the invention. The entire contents of the patents and other publications referred to in this specification are incorporated herein by reference.

25

PARTS LIST

201	compensation film
203	plane of the film
301	index ellipsoid approximating the state of liquid crystal cell 305
305	Liquid Crystal cell
307	compensation film
309	index ellipsoid representing the compensation film 307
311	obliquely propagating rays
401	dispersion curve of typical liquid crystal
403	dispersion curve for compensation film with proper dispersion
407	dispersion curve for compensation film with reverse dispersion
501	compensation film
503	polymer layer
505	substrate
507	additional layer
511	compensation film
513	compensation film
601	liquid crystal display
603	compensation film
605	liquid crystal cell
607	polarizer
609	polarizer
611	compensation film
613	liquid crystal display
701	dispersion curve (birefringence) of the example polymer layer A
703	dispersion curve (birefringence) of the example polymer layer B
801	dispersion curve (phase retardation) of the exemplary compensation film
901	dispersion curve (phase retardation) of the comparative example film
ϕ	angle of incidence of ray
n_x	index of refraction in x direction
n_y	index of refraction in y direction
n_z	index of refraction in z direction

n_{xlc}	index of refraction in x direction of liquid crystal cell
$n_{y lc}$	index of refraction in y direction of liquid crystal cell
$n_{z lc}$	index of refraction in z direction of liquid crystal cell
n_o	ordinary index of refraction
n_e	extraordinary index of refraction
$n_{o lc}$	ordinary index of refraction of liquid crystal
$n_{e lc}$	ordinary index of refraction of liquid crystal
Δn_{lc}	birefringence of liquid crystal
Δn_{th}	out-of-plane birefringence
d	thickness of the film
d_{lc}	thickness of the liquid crystal cell
R_{in}	in-plane phase retardation
R_{th}	out-of-plane phase retardation
R_{th-lc}	out-of-plane phase retardation of the liquid crystal cell
λ	wavelength

CLAIMS:

1. An optically transparent compensation film for Liquid Crystal Displays comprising an amorphous polymer layer disposed on a substrate wherein
5 said amorphous polymer layer has an out-of-plane birefringence $\Delta n_{th}(\lambda)$ at a wavelength λ that satisfies both conditions (1) and (2):

Condition (1)

$\Delta n_{th}(\lambda)$ is more negative than -0.005 throughout the range of $370\text{nm} < \lambda < 700\text{nm}$;

Condition (2)

- 10 $\Delta n_{th}(\lambda_1)$ is more negative than $\Delta n_{th}(\lambda_2)$, for $\lambda_1 < \lambda_2$ throughout the range of $370\text{nm} < \lambda < 700\text{nm}$.

2. An optically transparent compensation film according to claim 1 wherein, the out-of-plane phase retardation, R_{th} , of said compensation film
15 satisfies condition (3):

Condition (3)

R_{th} at λ_1 is more negative than R_{th} at λ_2 , for $\lambda_1 < \lambda_2$ throughout the range of $400\text{nm} < \lambda < 700\text{nm}$.

- 20 3. An optically transparent compensation film according to claim 2 wherein, said amorphous polymer layer is contiguous to the substrate.

4. An optically transparent compensation film according to claim 2 wherein, said amorphous polymer layer is contiguous to a barrier layer, which is
25 contiguous to the substrate.

5. An optically transparent compensation film according to claim 2 wherein, said substrate is optically uniaxial.

6. An optically transparent compensation film according to claim 5 wherein, said uniaxial substrate is negative-C plate and the R_{th} (at $\lambda = 550\text{nm}$ wavelength) is between -20 to -250nm .

5 7. An optically transparent compensation film according to claim 2 wherein, said substrate is optically biaxial.

8. An optically transparent compensation film according to claim 2 wherein, an additional layer is disposed on said amorphous polymer layer.

10

9. An optically transparent compensation film according to claim 8 wherein, said additional layer has an anti-reflection function.

10. An optically transparent compensation film according to claim 8
15 wherein, said additional layer has an anti-glare function.

11. An optically transparent compensation film according to claim 8 wherein, said additional layer is a hard-coat layer.

20 12. An optically transparent compensation film according to claim 2 wherein, said substrate is optically isotropic.

13. An optically transparent compensation film according to claim 2 wherein, said amorphous polymer layer has a thickness of from 0.1 to $30\mu\text{m}$.

25

14. An optically transparent compensation film according to claim 13 wherein, said amorphous polymer layer has a thickness of from 1 to $10\mu\text{m}$.

15 15. An optically transparent compensation film according to claim 2 wherein, said amorphous polymer layer comprises a polymer containing in the backbone a non-visible chromophore group and has a T_g above 180°C .

16. An optically transparent compensation film according to claim 15 wherein, said non-visible chromophore group includes a carbonyl, amide, ester, carbonate, phenyl, naphthyl, biphenyl, bisphenol, or thiophene group.

5 17. An optically transparent compensation film of claim 15 wherein, said amorphous polymer layer comprises a polymer containing in the backbone a non-visible chromophore group that does not contain a chromophore off of the backbone.

10 18. An optically transparent compensation film according to claim 2 wherein, said amorphous polymer layer comprises a polymer with Tg above 180°C.

15 19. An optically transparent compensation film according to claim 17 wherein, said amorphous polymer layer comprises a polymer containing in the backbone a vinyl, carbonyl, amide, ester, carbonate, aromatic, sulfone, or azo group.

20 20. An optically transparent compensation film according to claim 2 wherein, said amorphous polymer layer comprises 1)poly(4,4'-hexafluoroisopropylidene-bisphenol) terephthalate-co-isophthalate, 2)poly(4,4'-hexahydro-4,7-methanoindan-5-ylidene bisphenol) terephthalate, 3) poly(4,4'-isopropylidene-2,2'6,6'-tetrachlorobisphenol) terephthalate-co-isophthalate, 4) poly(4,4'-hexafluoroisopropylidene)-bisphenol-co-(2-norbornylidene)-bisphenol terephthalate, 5) poly(4,4'-hexahydro-4,7-methanoindan-5-ylidene)-bisphenol-co-
25 (4,4'-isopropylidene-2,2',6,6'-tetrabromo)-bisphenol terephthalate, or 6) poly(4,4'-isopropylidene-bisphenol-co- 4,4'-(2-norbornylidene) bisphenol) terephthalate-co-isophthalate or copolymers of any of the foregoing.

21. An optically transparent compensation film according to claim 2 wherein, said amorphous polymer layer comprises poly(4,4'-hexafluoroisopropylidene-bisphenol-co- 4,4'-(2-norbornylidene) bisphenol) terephthalate-co-isophthalate or copolymers thereof.

22. An optically transparent compensation film according to claim 2 wherein, said substrate comprises triacetylcellulose (TAC), polycarbonate, cyclic polyolefin or polyarylate containing fluorene groups.

23. An optically transparent compensation film according to claim 2 wherein, more than one layer of said amorphous polymer are disposed on said substrate.

24. An optically transparent compensation film according to claim 2 wherein, the thickness of said compensation film is less than 115 μ m.

25. An optically transparent compensation film according to claim 24 wherein, the thickness of said compensation film is between 40 to 100 μ m.

26. A liquid crystal display comprising a liquid crystal cell, a pair of crossed polarizers located one on each side of the cell, and at least one optical compensation film according to claim 2.

27. A liquid crystal display according to claim 26 wherein, said liquid crystal cell is a Vertically Aligned liquid crystal cell or Twisted Nematic liquid crystal cell.

28. A process for forming an optically transparent compensation film for an liquid crystal displays comprising coating an layer of amorphous polymer in a solvent onto a substrate wherein, said amorphous polymer layer satisfies the both conditions (1) and (2):

Condition (1)

$\Delta n_{th}(\lambda)$ is more negative than -0.005 throughout the range of $370\text{nm} < \lambda < 700\text{nm}$;

Condition (2)

5 $\Delta n_{th}(\lambda_1)$ is more negative than $\Delta n_{th}(\lambda_2)$, for $\lambda_1 < \lambda_2$ throughout the range of $370\text{nm} < \lambda < 700\text{nm}$,

and said amorphous polymer layer comprises selected polymeric materials and said layer has sufficient thickness so that the out-of-plane retardation R_{th} (at $\lambda = 550\text{nm}$ wavelength) of said amorphous polymer layers is more negative than

10 -5nm.

29. An optical compensation film for Liquid Crystal Displays comprising an amorphous polymer layer disposed on a polymeric substrate wherein said amorphous polymer layer has an out-of-plane birefringence $\Delta n_{th}(\lambda)$ at a wavelength λ that satisfies both conditions (1) and (2):

Condition (1)

$\Delta n_{th}(\lambda)$ is more negative than -0.005 throughout the range of $370\text{nm} < \lambda < 700\text{nm}$;

Condition (2)

20 $\Delta n_{th}(\lambda_1)$ is more negative than $\Delta n_{th}(\lambda_2)$, for $\lambda_1 < \lambda_2$ throughout the range of $370\text{nm} < \lambda < 700\text{nm}$.

30. An optical compensation film according to claim 29 wherein, said polymeric substrate comprises triacetylcellulose (TAC), polycarbonate, cyclic polyolefin or polyarylate containing fluorene groups.

25

31. An optical compensation film according to claim 29 wherein, said amorphous polymer layer comprises 1) poly(4,4'-hexafluoroisopropylidene-bisphenol) terephthalate-co-isophthalate, 2) poly(4,4'-hexahydro-4,7-methanoindan-5-ylidene bisphenol) terephthalate, 3) poly(4,4'-isopropylidene-30 2,2'6,6'-tetrachlorobisphenol) terephthalate-co-isophthalate, 4) poly(4,4'-hexafluoroisopropylidene)-bisphenol-co-(2-norbornylidene)-bisphenol

terephthalate, 5) poly(4,4'-hexahydro-4,7-methanoindan-5-ylidene)-bisphenol-co-(4,4'-isopropylidene-2,2',6,6'-tetrabromo)-bisphenol terephthalate, or 6) poly(4,4'-isopropylidene-bisphenol-co- 4,4'-(2-norbornylidene) bisphenol) terephthalate-co-isophthalate or copolymers of any of the foregoing.

5

32. An optical compensation film according to claim 29 wherein, said amorphous polymer layer comprises poly(4,4'-hexafluoroisopropylidene-bisphenol-co- 4,4'-(2-norbornylidene) bisphenol) terephthalate-co-isophthalate or copolymers thereof.

10

33. An optical compensation film according to claim 29 wherein, said amorphous polymer layer comprises a polymer with Tg above 180°C.

34. An optical compensation film according to claim 29 wherein, said
15 amorphous polymer layer comprises a polymer containing in the backbone a non-visible chromophore group and has a Tg above 180°C.

35. A liquid crystal display comprising a liquid crystal cell, a pair of
crossed polarizers located one on each side of the cell, and at least one optical
20 compensation film according to claim 29.

36. A liquid crystal display according to claim 35 wherein, said liquid
crystal cell is a Vertically Aligned liquid crystal cell or Twisted Nematic liquid
crystal cell.

25

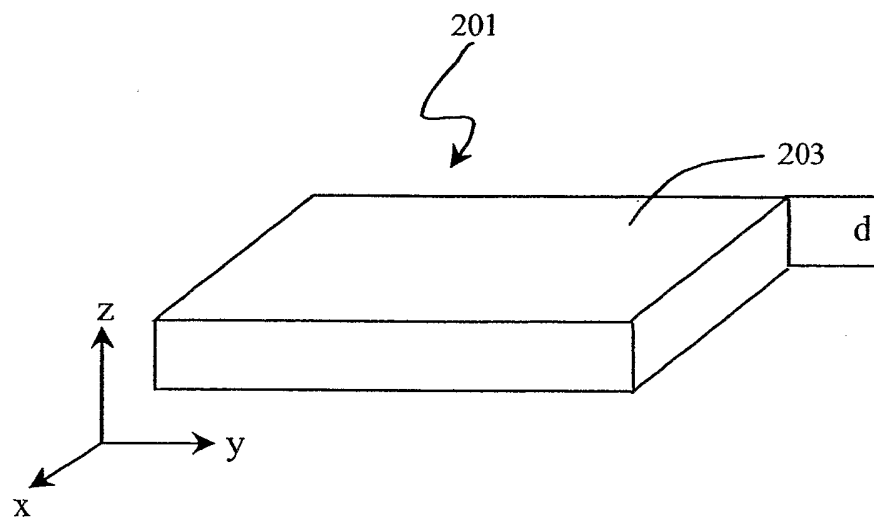


FIG. 1

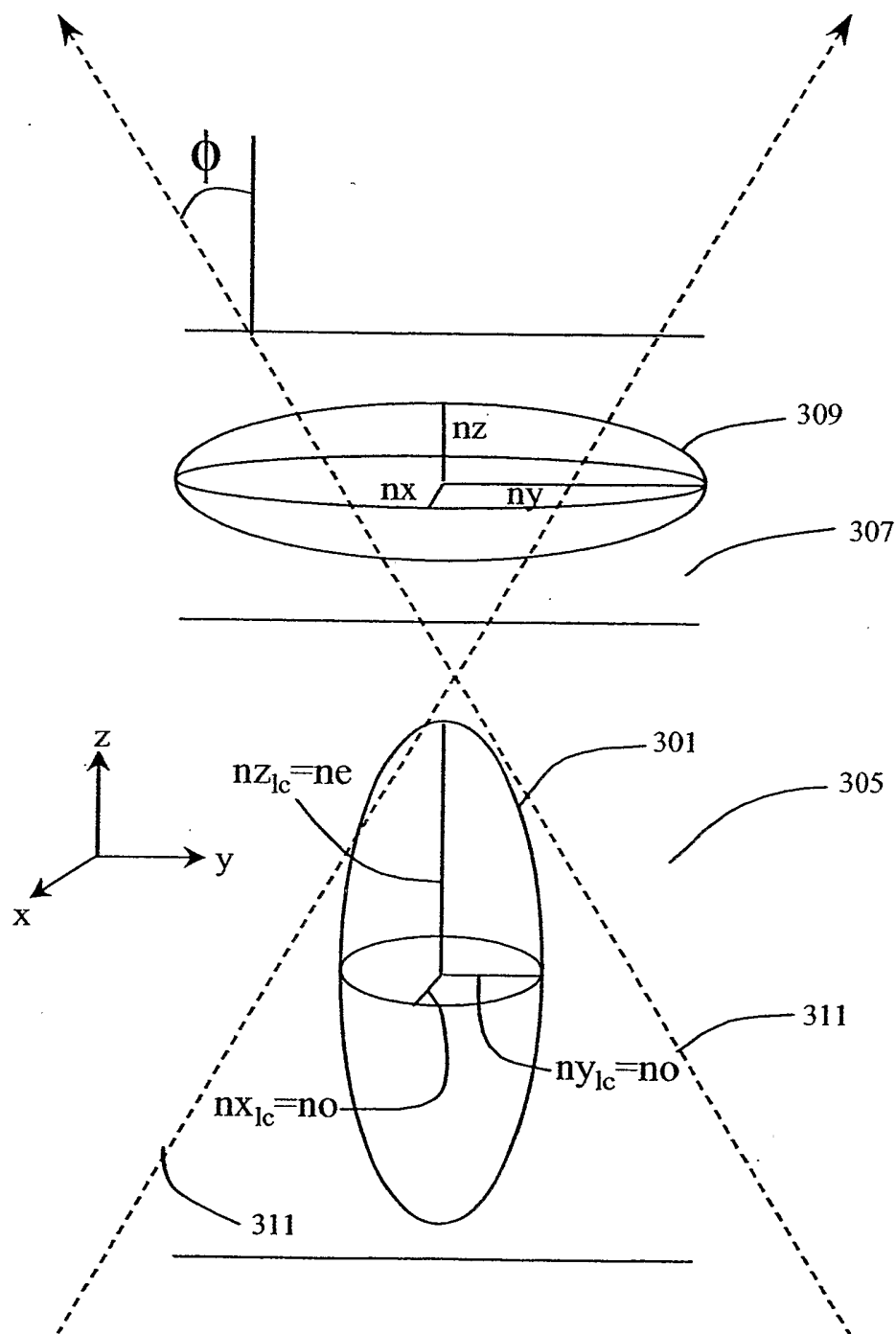


FIG. 2

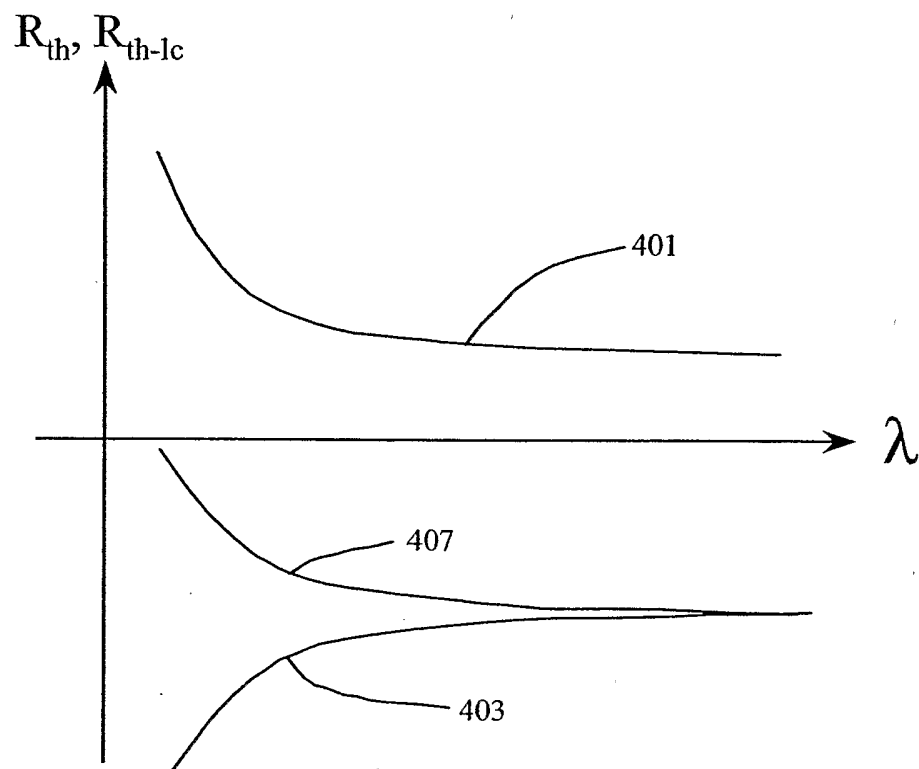


FIG. 3

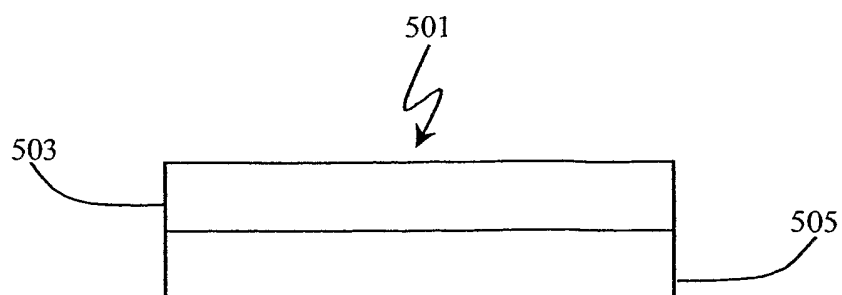


FIG. 4A

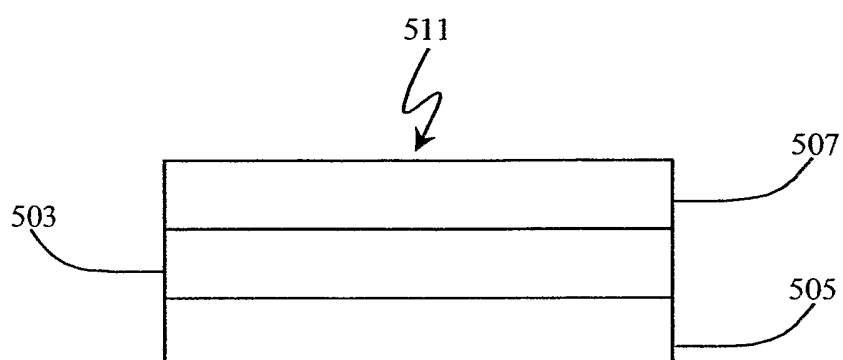


FIG. 4B

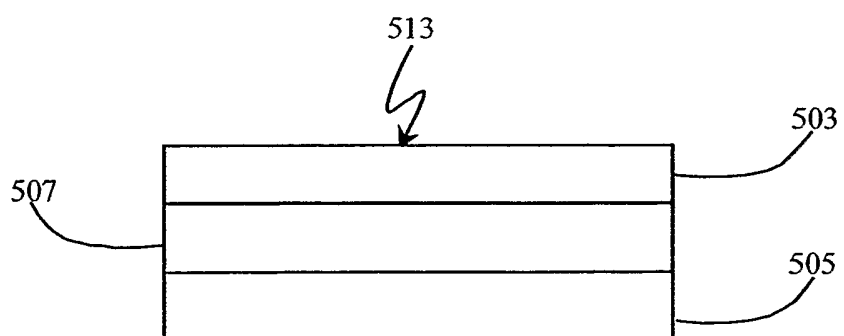


FIG. 4C

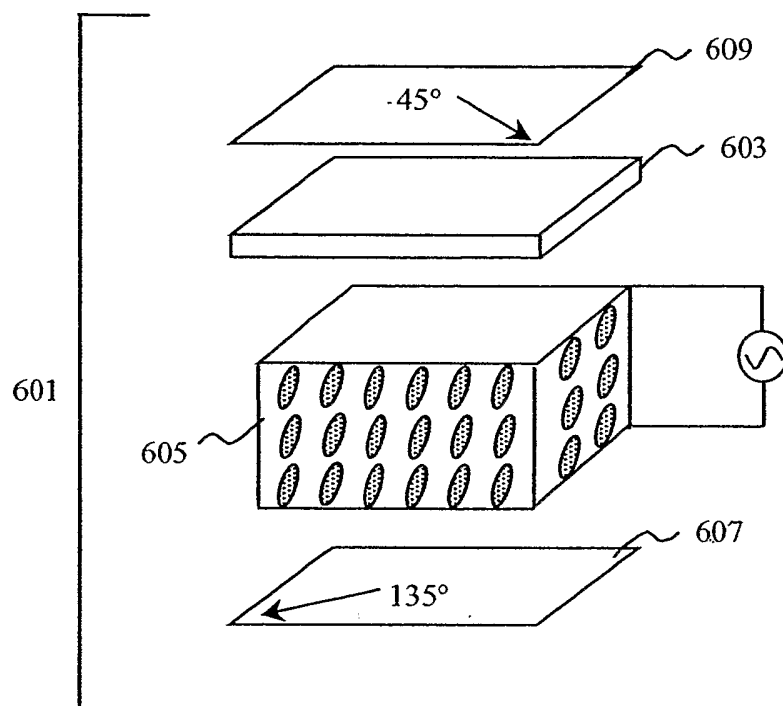


FIG. 5A

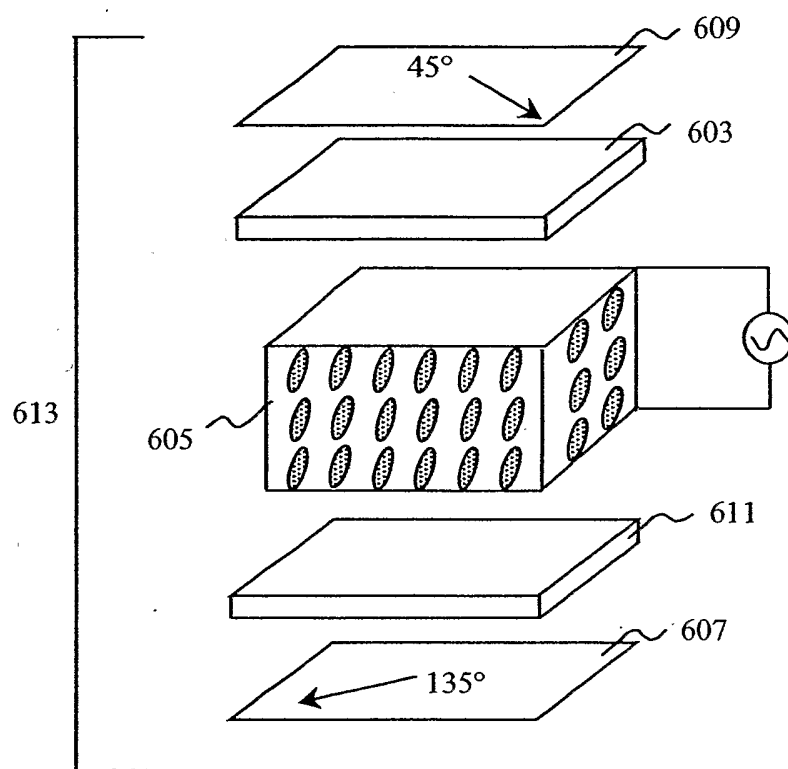


FIG. 5B

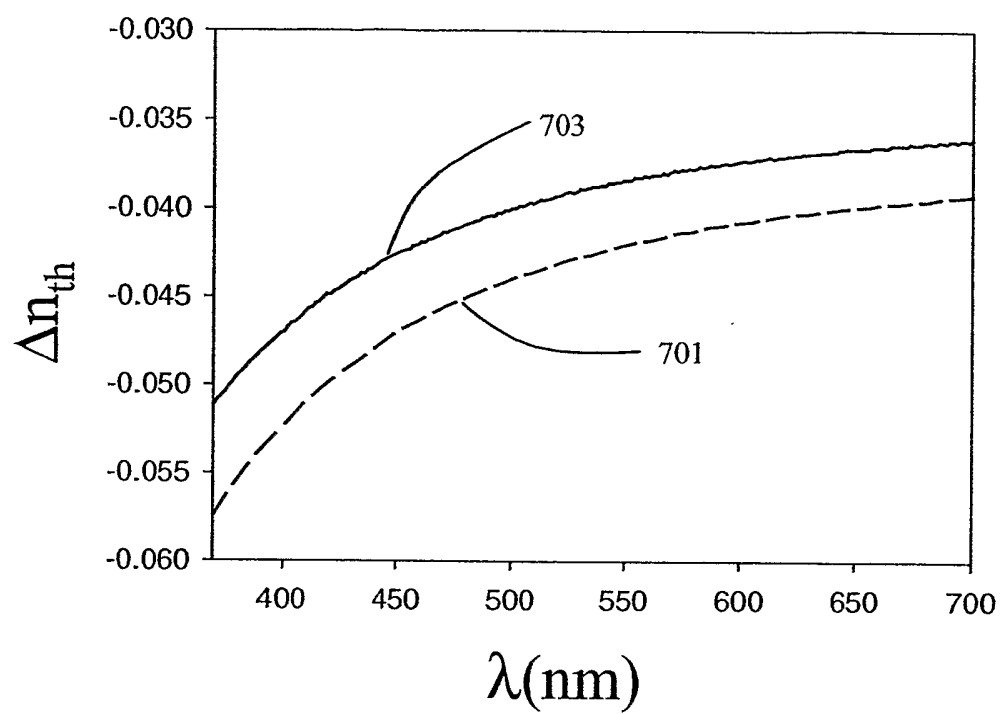


FIG.6

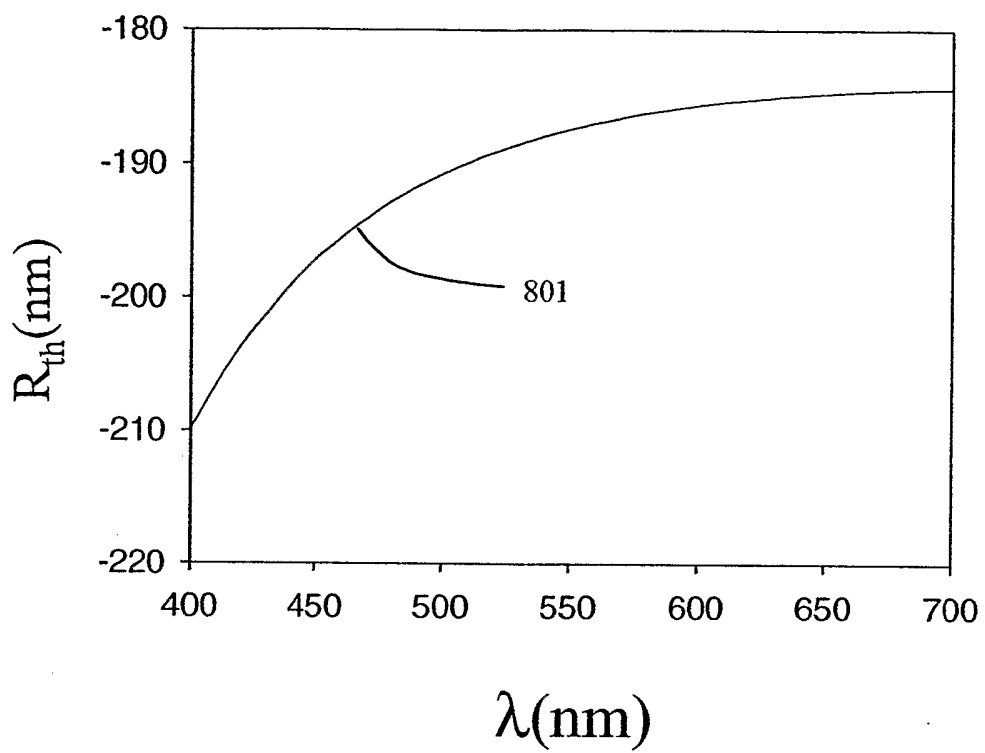


FIG.7

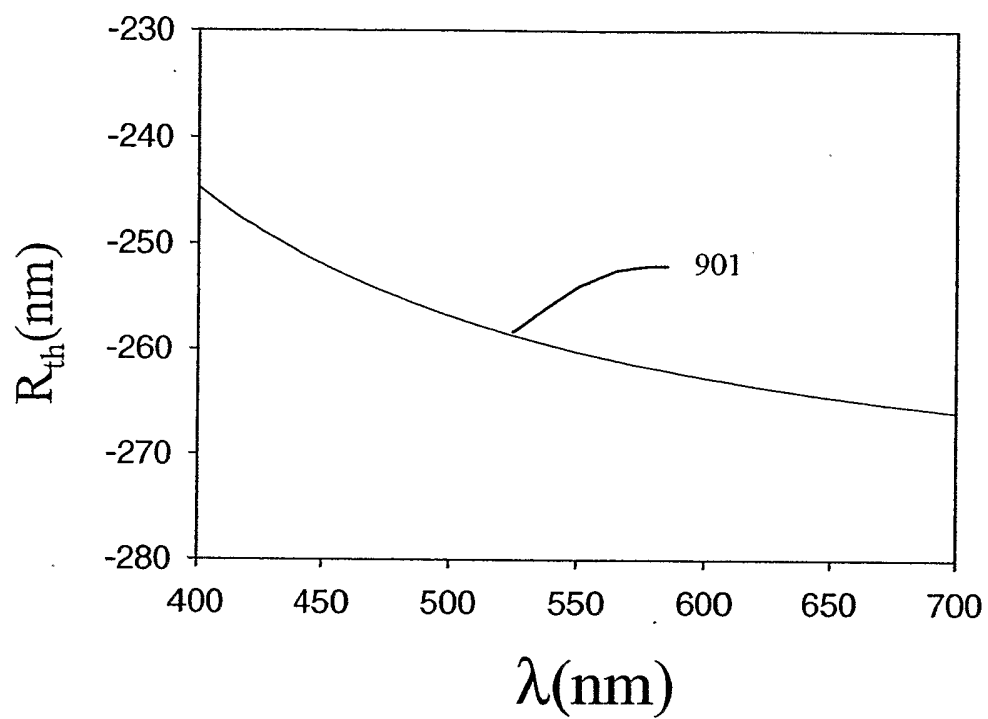


FIG. 8

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US2004/013676

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 G02F1/13363 C08G64/04

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 G02F C08G G02B

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

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 6 238 753 B1 (SAHOUBANI HASSAN ET AL) 29 May 2001 (2001-05-29) column 1, line 10 - line 27 column 2, line 15 - column 3, line 47 column 10, line 41 - column 11, line 56; figures 1,2 column 14, line 5 - line 29 column 21, line 58 - column 22, line 67; table 4 column 29, line 25 - column 30, line 52; tables 9,10 column 8, line 32 - line 52	1-8, 11-16, 18,19, 22-30, 33-36
Y	-/--	9,10



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
- *E* earlier document 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.
- *G* document member of the same patent family

Date of the actual completion of the international search

27 September 2004

Date of mailing of the international search report

22/10/2004

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Cossu, A

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US2004/013676

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	-& US 2002/192445 A1 (AYUKAWA HIROSHI ET AL) 19 December 2002 (2002-12-19) paragraph '0024! - paragraph '0029! -----	15,18, 33,34
X	US 5 480 964 A (CHENG STEPHEN Z D ET AL) 2 January 1996 (1996-01-02) column 4, line 10 - line 42 column 5, line 40 - column 23, line 45 column 26, line 25 - line 45; tables I,IV column 54, line 1 - line 25; table VI column 65, line 55 - column 67, line 67; figures 2,4	1-4,8, 15-19, 23,26,28
X	-& FUMING L ET AL: "Polyimide films as negative birefringent compensators for normally white twisted nematic liquid crystal displays" POLYMER, ELSEVIER SCIENCE PUBLISHERS B.V, GB, vol. 37, no. 23, 1 November 1996 (1996-11-01), pages 5321-5325, XP004069525 ISSN: 0032-3861 page 5322, right-hand column - page 5323, right-hand column; figures 2-5; table 1 -----	1-4,8, 15-19, 23,26,28
Y	US 2003/058393 A1 (KOZAKI SHUICHI ET AL) 27 March 2003 (2003-03-27) paragraph '0240! - paragraph '0242! paragraph '0281! - paragraph '0301! paragraph '0310! - paragraph '0316! paragraph '0335! - paragraph '0343!; figures 55,56,62,63 -----	9,10
X	EP 0 544 013 A (FUJIMORI KOGYO CO ; SEIKO EPSON CORP (JP)) 2 June 1993 (1993-06-02) page 3, line 24 - line 28 page 4, line 30 - page 6, line 38 page 8, line 30 - page 9, line 40; figure 1 claim 3 -----	1-3,8, 12,22, 24, 26-30, 35,36
A	EP 1 197 768 A (TEIJIN LTD) 17 April 2002 (2002-04-17) paragraph '0020! - paragraph '0052! paragraph '0057! - paragraphs '0073! - '0086! paragraph '0096! - paragraph '0101!; table 3 -----	20,21, 31,32
A	US 6 380 996 B1 (ITO YOJI ET AL) 30 April 2002 (2002-04-30) column 35, line 27 - column 37, line 20 -----	5-7

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US2004/013676

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 6238753	B1	29-05-2001	US 6074709 A	13-06-2000
			US 5750641 A	12-05-1998
			US 2002012758 A1	31-01-2002
			AU 7369996 A	09-12-1997
			CN 1219247 A	09-06-1999
			EP 0900406 A1	10-03-1999
			JP 2000511296 T	29-08-2000
			KR 2000015781 A	15-03-2000
			US 5969088 A	19-10-1999
			WO 9744704 A1	27-11-1997
US 2002192445	A1	19-12-2002	WO 02090452 A2	14-11-2002
US 5480964	A	02-01-1996	US 5344916 A	06-09-1994
			US 5580950 A	03-12-1996
			AU 6711694 A	08-11-1994
			CA 2160908 A1	27-10-1994
			DE 69424937 D1	20-07-2000
			DE 69424937 T2	11-01-2001
			EP 0698052 A1	28-02-1996
			WO 9424191 A1	27-10-1994
			JP 8511812 T	10-12-1996
US 2003058393	A1	27-03-2003	EP 0899605 A2	03-03-1999
			JP 3526533 B2	17-05-2004
			JP 11133413 A	21-05-1999
			JP 2004004950 A	08-01-2004
			JP 2004004951 A	08-01-2004
			US 6512561 B1	28-01-2003
			JP 2000019518 A	21-01-2000
			JP 2003307735 A	31-10-2003
EP 0544013	A	02-06-1993	DE 69209008 D1	18-04-1996
			DE 69209008 T2	05-09-1996
			EP 0544013 A1	02-06-1993
			US 5396355 A	07-03-1995
			WO 9222835 A1	23-12-1992
			JP 3313406 B2	12-08-2002
			JP 5196816 A	06-08-1993
EP 1197768	A	17-04-2002	EP 1197768 A1	17-04-2002
			CN 1383495 T	04-12-2002
			WO 0181959 A1	01-11-2001
			JP 2002014234 A	18-01-2002
			TW 533323 B	21-05-2003
			US 2003086027 A1	08-05-2003
US 6380996	B1	30-04-2002	JP 2001027706 A	30-01-2001
			JP 2000304931 A	02-11-2000
			EP 0928984 A2	14-07-1999
			JP 11352328 A	24-12-1999
			AU 2326200 A	04-09-2000
			EP 1156349 A1	21-11-2001
			WO 0049430 A1	24-08-2000