

(19) World Intellectual Property Organization
International Bureau



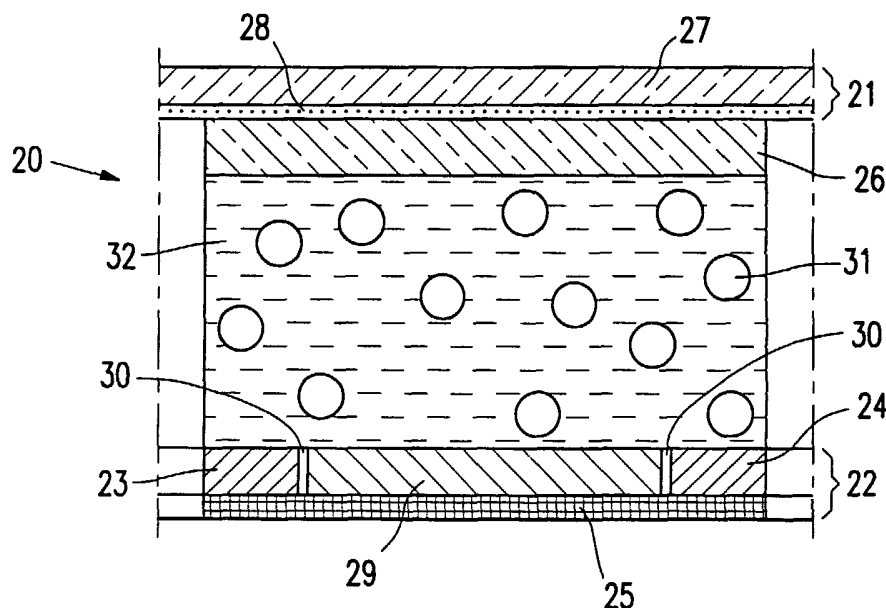
(43) International Publication Date
30 January 2003 (30.01.2003)

PCT

(10) International Publication Number
WO 03/009059 A1

- (51) International Patent Classification⁷: **G02F 1/167**
- (21) International Application Number: PCT/US02/22812
- (22) International Filing Date: 16 July 2002 (16.07.2002)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:
60/306,312 17 July 2001 (17.07.2001) US
- (71) Applicant: **SIPIX IMAGING, INC.** [US/US]; 1075 Montague Expressway, Milpitas, CA 95035 (US).
- (72) Inventors: **LIANG, Rong-Chang**; 20142 Pacifica Drive, Cupertino, CA 95014 (US). **CHUNG, Jerry**; 100 Montelena Court, Mountain View, CA 94040 (US). **CHEN, David**; 8312 Ridglea Avenue, Buena Park, CA 90621 (US).
- (74) Agent: **CHOW, Ping, Y.**; Heller Ehrman White & McAuliffe LLP, 275 Middlefield Road, Menlo Park, CA 94025-3506 (US).
- (81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, 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, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW.
- (84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, 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, IE, IT, LU, MC, NL, PT, SE, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).
- 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.

(54) Title: IN-PLANE SWITCHING ELECTROPHORETIC DISPLAY



(57) Abstract: The present invention relates to an improved EPD which comprises the in plane switching mode. More specifically, the EPD of the present invention comprises isolated cells formed from microcups of well defined size, shape and aspect ratio and the movement of the particles in the cells is controlled by the in-plane switching mode. The EPD of the present invention may be produced in a continuous manufacturing process, and the display gives improved color saturation.

IN-PLANE SWITCHING ELECTROPHORETIC DISPLAY

BACKGROUND

The electrophoretic display (EPD) is a non-emissive device based on the electrophoresis phenomenon influencing charged pigment particles suspended in a colored dielectric solvent. This general type of display was first proposed in 1969. An EPD typically comprises a pair of opposed, spaced-apart plate-like electrodes, with spacers predetermining a certain distance between the electrodes. At least one of the electrodes, typically on the viewing side, is transparent. For the passive type of EPDs, row and column electrodes on the top (the viewing side) and bottom plates respectively are needed to drive the displays. In contrast, an array of thin film transistors (TFT) on the bottom plate and a common, non-patterned transparent conductor plate on the top viewing substrate are required for the active type EPDs. An electrophoretic fluid composed of a colored dielectric solvent and charged pigment particles dispersed therein is enclosed between the two electrodes.

When a voltage difference is imposed between the two electrodes, the pigment particles migrate by attraction to the plate of polarity opposite that of the pigment particles. Thus, the color showing at the transparent plate, determined by selectively charging the plates, can be either the color of the solvent or the color of the pigment particles. Reversal of plate polarity will cause the particles to migrate back to the opposite plate, thereby reversing the color. Intermediate color density (or shades of gray) due to intermediate pigment density at the transparent plate may be obtained by controlling the plate charge through a range of voltages. No backlight is needed in this type of reflective EPD displays.

A transmissive EPD is disclosed in US Patent No. 6,184,856 in which a backlight, color filters, and substrates with two transparent electrodes are used. The electrophoretic cells serve as a light valve. In the collected state, the particles are positioned to minimize the coverage of the horizontal area of the cell and allow the backlight to pass through the cell. In the distributed state, the particles are positioned to cover the horizontal area of the pixel and scatter or absorb the backlight. However, the backlight and color filter used in this device consume a great deal of power and are not desirable for hand-held devices such as PDAs (personal digital assistants) and e-books.

Besides the normal top/bottom electrode switching mode of EPDs, reflective "in-plane" switching EPDs have been disclosed in E. Kishi, et al., "5.1: development of In-Plane EPD", Canon Research Center, SID 00 Digest, pages 24-27 (2000) and Sally A. Swanson, et al., "5.2: High Performance Electrophoretic Displays", IBM Almaden Research Center, SID 00 Digest, pages 29-31, (2000). However, only monochrome in-plane switching EPDs are disclosed in these references. To prepare a multicolor display, either color filters or isolated color pixels or cell structures are needed for color separation and rendition. Color filter is typically expensive and not power-efficient. On the other hand, the preparation of isolated pixels or cells for color separation and rendering in the in-plane switching mode has not been taught previously.

EPDs of different pixel or cell structures have been reported in prior art, for example, the partition-type EPD (M.A. Hopper and V. Novotny, IEEE Trans. Electr. Dev., Vol ED 26, No. 8, pp 1148-1152 (1979)) and the microencapsulated EPD (US Patent Nos. 5,961,804 and 5,930,026), and each of these has its own problems as noted below.

In a partition-type EPD, there are partitions between the two electrodes for dividing the space into smaller cells in order to prevent undesired movements of the particles such as sedimentation. However, difficulties are encountered in the formation of the partitions, the process of filling the display with the fluid, enclosing the fluid in the display, and keeping the suspensions of different colors separated from each other.

The microencapsulated EPD has a substantially two dimensional arrangement of microcapsules each having therein an electrophoretic composition of a dielectric fluid and a dispersion of charged pigment particles that visually contrast with the dielectric solvent. The microcapsules are typically prepared in an aqueous solution and, to achieve a useful contrast ratio, their mean particle size is relatively large (50-150 microns). The large microcapsule size results in a poor scratch resistance and a slow response time for a given voltage because a large gap between the two opposite electrodes is required for large capsules. Also, the hydrophilic shell of microcapsules prepared in an aqueous solution typically results in sensitivity to high moisture and temperature conditions. If the microcapsules are embedded in a large quantity of a polymer matrix to obviate these shortcomings, the use of the matrix results in an even slower response time and/or a lower contrast ratio. To improve the switching rate, a charge-controlling agent is often needed in this type of EPDs. However, the

microencapsulation process in an aqueous solution imposes a limitation on the type of charge-controlling agents that can be used. Other drawbacks associated with the microcapsule system include poor resolution and poor addressability for color applications.

5 An improved EPD technology was recently disclosed in co-pending applications, US Serial Number 09/518,488, filed on March 3, 2000 (corresponding to WO01/67170), US Serial Number 09/759,212, filed on January 11, 2001, US Serial Number 09/606,654, filed on June 28, 2000 (corresponding to WO02/01281) and US Serial
10 Number 09/784,972, filed on February 15, 2001, all of which are incorporated herein by reference. The improved EPD comprises closed isolated cells formed from microcups of well-defined shape, size and aspect ratio and filled with charged pigment particles dispersed in a dielectric solvent. The electrophoretic fluid is isolated and sealed in each microcup.

 The microcup structure, in fact, enables a format flexible, efficient roll-to-roll
15 continuous manufacturing process for the preparation of EPDs. The displays can be prepared on a continuous web of conductor film such as ITO/PET by, for example, (1) coating a radiation curable composition onto the ITO/PET film, (2) making the microcup structure by a microembossing or photolithographic method, (3) filling the electrophoretic fluid and sealing the microcups, (4) laminating the sealed microcups
20 with the other conductor film, and (5) slicing and cutting the display to a desirable size or format for assembling.

 One advantage of this EPD design is that the microcup wall is in fact a built-in spacer to keep the top and bottom substrates apart at a fixed distance. The mechanical properties and structural integrity of microcup displays are significantly better than any
25 prior art displays including those manufactured by using spacer particles. In addition, displays involving microcups have desirable mechanical properties including reliable display performance when the display is bent, rolled, or under compression pressure from, for example, a touch screen application. The use of the microcup technology also eliminates the need of an edge seal adhesive which would limit and predefine the size of
30 the display panel and confine the display fluid inside a predefined area. The display fluid in a conventional display prepared by the edge sealing adhesive method will leak out completely if the display is cut in any way, or if a hole is drilled through the display. The damaged display will be no longer functional. In contrast, the display fluid in the

display prepared by the microcup technology is enclosed and isolated in each cell. The microcup display may be cut to almost any dimension without the risk of damaging the display performance due to the loss of display fluids in the active areas. In other words, the microcup structure enables a format flexible display manufacturing process, wherein
5 the process produces a continuous output of displays in a large sheet format which can be sliced and diced to any desired format. The isolated microcup or cell structure is particularly important when cells are filled with fluids of different specific properties such as colors and switching rates. Without the microcup structure, it will be very difficult to prevent the fluids in adjacent areas from intermixing, or being subject to
10 cross-talk during operation.

Multi-color displays may thus be manufactured by using a spatially adjacent array of small pixels formed of microcups filled with dyes of different colors (*e.g.*, red, green or blue). However there is a major deficiency in this type of system with the traditional top/bottom electrode switching mode. The white light reflected from the
15 “turned-off” colored pixels greatly reduces the color saturation of the “turned-on” colors. More details in this regard are given in the following “Detailed Description” section.

While this deficiency may be remedied by an overlaid shutter device such as a polymer dispersed liquid crystal to switch each pixel to the black color, the disadvantage
20 of this approach is the high cost of the overlaid device and the complicated driving circuit design.

Thus, there is still a need for an EPD with improved properties that can also be prepared in an efficient manner.

SUMMARY OF THE INVENTION

25 The present invention relates to an improved EPD which comprises the in plane switching mode for image formation. More specifically, the EPD of the present invention comprises isolated cells formed from microcups of well defined size, shape and aspect ratio and the movement of the particles in the cells is controlled by the in-plane switching mode. The EPDs of the present invention may be produced in a
30 continuous roll-to-roll manufacturing process, and the resultant displays have improved color saturation and contrast ratio.

BRIEF DESCRIPTION OF THE DRAWINGS

It is noted that all figures are shown as schematic and are not to scale.

Figure 1 illustrates the common deficiency of the traditional EPD with only the top/bottom switching mode.

5 Figure 2 illustrates a typical electrophoretic cell of the present invention and the general locations of the in-plane switching electrodes.

Figures 3A and 3B illustrate the monochrome display of the present invention.

Figures 4A-4D illustrate the various multiple color scenarios of the present invention.

10 Figures 5A and 5B illustrate the manufacture of microcups involving imagewise photolithographic exposure through a photomask.

DETAILED DESCRIPTION OF THE INVENTION

Unless defined otherwise in this specification, all technical terms are used herein according to their conventional definitions as they are commonly used and understood
15 by those of ordinary skill in the art. The terms “cell”, “microcup”, “well-defined”, “aspect ratio”, and “imagewise exposure” are as defined in the co-pending applications identified above.

The term “isolated” refers to the electrophoretic cells which are individually sealed and the fluid in the cells may not be transferred from one cell to the other cells.

20 I. The Disadvantages of Electrophoretic Display with the Traditional Top/Bottom Switching

The EPD of Figure 1 has conventional top/bottom electrode switching mode. The cells are filled with a suspension in which white charged particles are dispersed in a colored (red, green and blue) dielectric solvent. All three cells in Figure 1 are shown
25 charged with a voltage difference between the top and bottom electrodes (not shown). In the green and blue cells, the white particles migrate to the top viewing electrode which is transparent, and as a result, the color of the particles (*i.e.*, white) is reflected to the viewer through the transparent conductor film in the two cells. In the red cell, the white particles migrate to the bottom of the cell, and the color of the solvent (*i.e.*, red) is
30 seen through the top transparent conductor film. In Figure 1, the white light reflected

from the green and blue cells dramatically reduces the saturation and contrast ratio of the red color.

In addition to the above mentioned problem, low solubility and poor fastness of the dyes in dielectric solvents of very low polarity, such as perfluoro and hydrocarbon solvents, have been a challenge for achieving high contrast ratio in the top/down type of EPDs.

II. Electrophoretic Display of the Present Invention

Figure 2 illustrates a typical electrophoretic cell of the present invention. The cell (20) comprises a top layer (21) and a bottom layer (22). The bottom layer has the in-plane switching electrodes (23) and (24) and the background layer (25). There is a common electrode (29) between the two in-plane electrodes separated by gaps (30). Alternatively, the bottom layer may have only one in-plane switching electrode, and one common electrode with a gap in between. Another alternative is where the background layer (25) is on top of the electrodes in the bottom layer (not shown). The in-plane electrode layer may also serve as the background layer and in this case the in-plane electrode(s) may be white or colored.

Typically, the cells of Figure 2 are filled with a dispersion of colored particles (31) in a clear dielectric solvent (32). The particles may be white, black or colored (*i.e.*, red, green or blue). The background layer (25) may be colorless, white, black or colored. The filled cells are subsequently sealed with a sealing layer (26). The top layer (21) having a transparent insulator layer (27) and preferably an adhesive layer (28) is then laminated over the sealed cells.

Preferably the microcup array is prepared in an up-side-down manner. In this scenario, the microcup array is prepared on the top transparent insulator substrate by either microembossing or photolithography as disclosed in the co-pending patent applications, US Serial Number 09/518,488, filed on March 3, 2000 (corresponding to WO01/67170), US Serial Number 09/759,212, filed on January 11, 2001, US Serial Number 09/606,654, filed on June 28, 2000 (corresponding to WO02/01281) and US Serial Number 09/784,972, filed on February 15, 2001, all of which are incorporated herein by reference. The microcups are filled with the electrophoretic fluid and sealed subsequently with a sealing layer. The bottom layer which contains the patterned electrodes and preferably an adhesive layer is then laminated over the sealed microcup.

The color background may be added by painting, printing, coating or laminating a color layer to the bottom electrode substrate.

One of the advantages of the in-plane switching mode is the possibility of making the microcups on a clear plastic insulator substrate. This eliminates the risk of breaking the brittle conductor electrode such as ITO/PET during the microembossing and other web handling steps. The patterned in-plane conductor film will only be used at the last step for lamination onto the filled and sealed microcups to complete the making of the display panel.

(1) Reflective Monochrome Display

In the cell as shown in Figure 3A, white particles are dispersed in a clear, colorless dielectric solvent. The background of all cells is of the same color (black, blue, cyan, red, magenta, etc.). When there is a voltage difference between the common (not shown) and the two in-plane switching electrodes (not shown), the white particles migrate to the sides of the cells, resulting in the color of the background being seen through the top transparent opening. When there is no voltage difference between the common and the two in-plane electrodes, the white particles are distributed in the dielectric solvent and as a result, the color of the particles (*i.e.*, white) is seen through the top transparent insulator layer.

Alternatively, as shown in Figure 3B, particles of the same color are dispersed in a clear, colorless dielectric solvent in all cells and the background of the cells is white. When there is a voltage difference between the common (not shown) and the two in-plane switching electrodes (not shown), the colored particles migrate to the sides of the cells, resulting in the color of the background (*i.e.*, white) being seen through the top transparent opening. When there is no voltage difference between the two in-plane electrodes, the colored particles are distributed in the dielectric solvent and as a result, the color of the particles is seen through the top transparent layer.

(2) Reflective Multiple Color Display

Figures 4A-4D illustrate the multiple color displays of the present invention.

In Figure 4A, the cells are filled with a colorless dielectric solvent with white charged particles dispersed therein, and have different background colors (*i.e.*, red, green or blue). When there is a voltage difference between the in-plane electrodes (not shown), the white particles migrate to either side of the cell, the color of the background

(*i.e.*, red, green or blue) is seen from the top transparent opening. When there is no voltage difference between the in-plane electrodes, the particles are distributed in the dielectric solvent resulting in the white color (*i.e.*, the color of the particles) being seen from the top transparent opening.

5 In Figure 4B, the cells are filled with a colorless dielectric solvent with black particles dispersed therein, and have different background colors (*i.e.*, red, green or blue). When there is a voltage difference between the in-plane electrodes (not shown), the particles migrate to either side of the cell, the color of the background (*i.e.*, red, green or blue) is seen from the top transparent opening. When there is no voltage
10 difference between the in-plane electrodes, the particles are distributed in the dielectric solvent, resulting in the black color (*i.e.*, the color of the particles) being seen from the top transparent opening.

 Figure 4C shows the cells filled with a colorless dielectric solvent with particles of different colors (*i.e.*, red, green or blue) dispersed therein. The background of the
15 cells is black. When there is a voltage difference between the in-plane electrodes (not shown), the colored charged particles migrate to either side of the cell, the color of the background (*i.e.*, black) is seen from the top transparent opening. When there is no voltage difference between the in-plane electrodes, the colored particles are distributed in the dielectric solvent, resulting in the color of the particles (*i.e.*, red, green or blue)
20 being seen from the top transparent opening. In this design, the black state is of high quality.

 In Figure 4D, the cells are filled with a colorless dielectric solvent with particles of different colors (red, green or blue) dispersed therein. The background of the cells is white. When there is a voltage difference between the in-plane electrodes (not shown),
25 the particles migrate to either side of the cell, the color of the background (*i.e.*, white) is seen from the top transparent opening, resulting in a high quality white state. When there is no voltage difference between the in-plane electrodes, the particles are distributed in the dielectric solvent, resulting in the color of the particles (*i.e.*, red, green or blue) being seen from the top transparent opening.

30 As shown in Figures 4A-4D, the in-plane switching mode allows the particles to move in the planar (left/right) direction, and different color combinations of particles, background, and fluid, wherein each is individually white, black, red, green or blue, can be used to generate various multi-color EPDs.

In addition, the particles in the dielectric solvent may be of mixed colors and the cells have the same background color.

In an alternative reflective display of the present invention, the top transparent viewing layer of the display may be colored or by adding a color filter. In this case, the cells are filled with an electrophoretic composition comprising white charged particles in a clear colorless or colored dielectric solvent and the background of the cells may be black. In a monochrome display, the transparent viewing layer on each pixel is of the same color (such as black, red, green, blue, yellow, cyan, magenta, etc.). In a multiple color display, the transparent viewing layers may be of different colors.

10 **III. Preparation of Microcup Array of the Present Invention**

The microcups generally may be manufactured by microembossing or photolithography as disclosed in US Patent Application Nos. USSN 09/518,488 filed March 3, 2000 (corresponding to WO01/67170) and USSN 09/784,972 filed on February 15, 2001.

15 **III(a) Preparation Of The Microcup Array by Microembossing** **Preparation of the Male Mold**

The male mold may be prepared by any appropriate method, such as a diamond turn process or a photoresist process followed by either etching or electroplating. A master template for the male mold may be manufactured by any appropriate method, such as electroplating. With electroplating, a glass base is sputtered with a thin layer (typically 3000 Å) of a seed metal such as chrome inconel. It is then coated with a layer of photoresist and exposed to radiation, such as ultraviolet (UV). A mask is placed between the UV and the layer of photoresist. The exposed areas of the photoresist become hardened. The unexposed areas are then removed by washing them with an appropriate solvent. The remaining hardened photoresist is dried and sputtered again with a thin layer of seed metal. The master is then ready for electroforming. A typical material used for electroforming is nickel cobalt. Alternatively, the master can be made of nickel by electroforming or electroless nickel deposition as described in "Continuous manufacturing of thin cover sheet optical media", SPIE Proc. Vol. 1663, pp. 324 (1992). The floor of the mold is typically between about 50 to 400 microns thick. The master can also be made using other microengineering techniques including e-beam writing, dry etching, chemical etching, laser writing or laser interference as described in

“Replication techniques for micro-optics”, SPIE Proc. Vol. 3099, pp. 76-82 (1997).

Alternatively, the mold can be made by photomachining using plastics, ceramics or metals.

5 The male mold thus prepared typically has protrusions between about 1 to 500 microns, preferably between about 2 to 100 microns, and most preferably about 4 to 50 microns. The male mold may be in the form of a belt, a roller, or a sheet. For continuous manufacturing, the belt type of mold is preferred. Prior to applying a UV curable resin composition, the mold may be treated with a mold release to aid in the demolding process.

10 Microcups may be formed either in a batchwise process or in a continuous roll-to-roll process as described in USSN 09/784,972 filed on February 15, 2001.

In the first step of the microembossing process, a UV curable resin is first coated on a substrate, preferably a transparent insulator, by any appropriate means, such as roller coating, die coating, slot coating, slit coating, doctor blade coating, and the like.
15 Suitable transparent insulator substrates include polyethylene terephthalate, polyethylene naphthate, polyaramid, polyimide, polycycloolefin, polysulfone, epoxy and their composites. The radiation curable material used is a thermoplastic or thermoset precursor, such as multifunctional acrylate or methacrylate, vinyl ether, epoxide and their oligomers, polymers and the like. Multifunctional acrylates and their
20 oligomers are the most preferred. A combination of a multifunctional epoxide and a multifunctional acrylate is also very useful to achieve desirable physico-mechanical properties. The UV curable resin may be degassed prior to dispensing and may optionally contain a solvent. The solvent, if present, readily evaporates.

The radiation curable material coated on the substrate is embossed by the male
25 mold under pressure. If the male mold is metallic and opaque, the plastic insulator is typically transparent to the actinic radiation used to cure the resin. Conversely, the male mold can be transparent and the plastic insulator can be opaque to the actinic radiation. The plastic insulator is preferably transparent because it is typically the viewing side. In this case, the electrodes can be opaque. Alternatively, the microembossing can be
30 performed on the substrate containing the electrodes.

After exposure to radiation, the radiation curable material becomes hardened. The male mode is then removed exposing the microcups formed.

III(b) Preparation of Microcup Array by Photolithography

The photolithographic process for preparation of the microcup array is shown in Figures 5A and 5B.

As shown in Figures 5A and 5B, the microcup array (50) may be prepared by exposure of a radiation curable material (51a), coated by any known methods onto an insulator substrate base (53), to UV light (or alternatively other forms of radiation, electron beams and the like) through a mask (56) to form walls (51b) corresponding to the image projected through the mask (56).

In the photomask (56) in Fig. 5A, the dark squares (54) represent the area opaque to the radiation employed, and the space (55) between the dark squares represents the radiation-transparent area. The UV radiates through the opening area (55) onto the radiation curable material (51a).

As shown in Figure 5B, the exposed areas (51b) become hardened and the unexposed areas (protected by the opaque area (54) of the mask (56)) are then removed by an appropriate solvent or developer to form the microcups (57). The solvent or developer is selected from those commonly used for dissolving or dispersing radiation curable materials such as methylethylketone, toluene, acetone, isopropanol or the like.

Alternatively, the exposure can be done by placing the photomask underneath the insulator substrate. In this case, the substrate must be transparent to the radiation wavelength used for exposure.

The openings of the microcups prepared according to the methods described above may be round, square, rectangular, hexagonal, or any other shape. The partition area between the openings is preferably kept small in order to achieve a high color saturation and contrast while maintaining desirable mechanical properties. Consequently the honeycomb-shaped opening is preferred over, for example, the circular opening.

For reflective electrophoretic displays, the dimension of each individual microcup may be in the range of about 10^2 to about $1 \times 10^6 \mu\text{m}^2$, preferably from about 10^3 to about $1 \times 10^5 \mu\text{m}^2$. The depth of the microcups is in the range of about 5 to about 200 microns, preferably from about 20 to about 100 microns. The opening to the total area ratio, total area being defined as that of one cup including walls measured from wall centers, is in the range of from about 0.2 to about 0.95, preferably from about 0.5 to about 0.9. The distances of the openings usually are in the range of from about 15 to

about 450 microns, preferably from about 25 to about 300 microns from edge to edge of the openings.

III(c) Sealing of the Microcups

After the microcups are filled with an electrophoretic fluid, they are sealed. The critical step of sealing of the microcups may be accomplished in a number of ways. A preferred approach is to disperse a UV curable composition into an electrophoretic fluid comprising charged pigment particles dispersed in a colored dielectric solvent. The suitable UV curable materials include acrylates, methacrylates, styrene, alpha-methylstyrene, butadiene, isoprene, allyacrylate, polyvalent acrylate or methacrylate, cyanoacrylates, polyvalent vinyl including vinylbenzene, vinylsilane, vinyl ether, polyvalent epoxide, polyvalent isocyanate, polyvalent allyl, and oligomers or polymers containing crosslinkable functional groups. The UV curable composition is immiscible with the dielectric solvent and has a specific gravity lower than that of the electrophoretic fluid, *i.e.*, the combination of the dielectric solvent and the pigment particles. The two components, UV curable composition and the electrophoretic fluid, are thoroughly blended in an in-line mixer and immediately coated onto the microcups with a precision coating mechanism such as Myrad bar, gravure, doctor blade, slot coating or slit coating. Excess fluid is removed by a wiper blade or a similar device. A small amount of a weak solvent or solvent mixture such as isopropanol or methanol may be used to clean the residual electrophoretic fluid on the top surface of the partition walls of the microcups. Volatile organic solvents may be used to control the viscosity and coverage of the electrophoretic fluid. The thus-filled microcups are then dried and the UV curable composition floats to the top of the electrophoretic fluid. The microcups may be sealed by curing the supernatant UV curable layer during or after it floats to the top. UV or other forms of radiation such as visible light, IR and electron beam may be used to cure the sealing layer and seal the microcups. Alternatively, heat or moisture may also be employed to cure the sealing layer and seal the microcups, if heat or moisture curable compositions are used.

A preferred group of dielectric solvents exhibiting desirable density and solubility discrimination against acrylate monomers and oligomers are halogenated hydrocarbons and their derivatives. Surfactants may be used to improve the adhesion and wetting at the interface between the electrophoretic fluid and the sealing materials.

Surfactants include the FC surfactants from 3M Company, Zonyl fluorosurfactants from DuPont, fluoroacrylates, fluoromethacrylates, fluoro-substituted long chain alcohols, perfluoro-substituted long chain carboxylic acids and their derivatives.

Alternatively, the electrophoretic fluid and the sealing precursor may be coated
5 sequentially into the microcups to prevent intermixing, if the sealing precursor is at least partially compatible with the dielectric solvent. Thus, the sealing of the microcups may be accomplished by overcoating a thin layer of sealing material which is hardenable by radiation, heat, moisture or interfacial reactions on the surface of the filled microcups. Volatile organic solvents may be used to adjust the viscosity and the thickness of the
10 coatings. When a volatile solvent is used in the overcoat, it is preferred that it is immiscible with the dielectric solvent to reduce the degree of intermixing between the sealing layer and the electrophoretic fluid. To further reduce the degree of intermixing, it is highly desirable that the specific gravity of the overcoating is significantly lower than that of the electrophoretic fluid. In the copending patent application, US Serial No.
15 09/874,391, filed June 4, 2001, thermoplastic elastomers have been disclosed as the preferred sealing material.

Examples of useful thermoplastic elastomers include ABA, and (AB)_n type of di-block, tri-block, and multi-block copolymers wherein A is styrene, α -methylstyrene, ethylene, propylene or norbornene; B is butadiene, isoprene, ethylene, propylene,
20 butylene, dimethylsiloxane or propylene sulfide; and A and B cannot be the same in the formula. The number, n, is ≥ 1 , preferably 1-10. Particularly useful are di-block or tri-block copolymers of styrene or α -methylstyrene such as SB (poly(styrene-b-butadiene)), SBS (poly(styrene-b-butadiene-b-styrene)), SIS (poly(styrene-b-isoprene-b-styrene)), SEBS (poly(styrene-b-ethylene/butylenes-b-styrene)) poly(styrene-b-dimethylsiloxane-
25 b-styrene), poly(α -methylstyrene-b-isoprene), poly(α -methylstyrene-b-isoprene-b- α -methylstyrene), poly(α -methylstyrene-b-propylene sulfide-b- α -methylstyrene), poly(α -methylstyrene-b-dimethylsiloxane-b- α -methylstyrene).

Alternatively, interfacial polymerization followed by UV curing has been found very beneficial to the sealing process. Intermixing between the electrophoretic layer and
30 the overcoat is significantly suppressed by the formation of a thin barrier layer at the interface by interfacial polymerization. The sealing is then completed by a post curing

step, preferably by UV radiation. The two-step overcoating process is particularly useful when the dye used is at least partially soluble in the thermoset precursor.

III(d) Lamination of the Microcups

5 The sealed microcups are then laminated with a top layer comprising a patterned in plane conductor film and preferably an adhesive layer. Suitable adhesive materials include acrylic and rubber types of pressure sensitive adhesives, UV curable adhesives containing for example, multifunctional acrylates, epoxides, or vinyl ethers, and moisture or heat curable adhesives such as epoxy, polyurethane, and cyanoacrylate.

10 The cells prepared from the methods of Sections III(a)-III(d) may be used in an up-side-down manner with the transparent viewing layer at the top and the layer with the in-plane electrodes at the bottom.

III(e) Alternative Methods

Alternatively, in the microembossing process, the UV curable resin is dispensed over the male mold by any appropriate means, such as coating, dipping, pouring and the like. The dispenser may be moving or stationary. A patterned in-plane conductor film on a plastic substrate such as polyethylene terephthalate, polyethylene naphthate, polyaramid, polyimide, polycycloolefin, polysulfone, epoxy and their composites is then overlaid on the UV curable resin. Pressure may be applied to ensure proper bonding between the resin and the plastic substrate and to control the thickness of the floor of the microcups. If the male mold is metallic and opaque, the plastic substrate is typically transparent to the actinic radiation used to cure the resin. Conversely, the male mold can be transparent and the plastic substrate can be opaque to the actinic radiation.

20 After exposure to UV radiation, the UV curable resin becomes hardened, and the male mode may then be removed. The microcup arrays formed are filled and sealed as described above. The sealed microcups are then laminated with a transparent insulator layer preferably using an adhesive.

Although less preferred, the photolithographic exposure may also be performed on the substrate having the in-plane electrodes. A radiation curable material is coated on the patterned conductor film. The microcups are formed by exposure of the radiation curable material to radiation through a photomask as shown in Figure 5 and described in Section III(b) above.

The microcups thus prepared are then filled and sealed as described above and laminated with a transparent insulator layer, preferably with an adhesive.

In any of the methods for the preparation of the microcups disclosed in this section, a substrate containing an array of thin film transistors (TFT) may be used as the
5 bottom in-plane electrode layer and in this case the TFT layer also provides an active driving mechanism.

IV. Preparation of the Suspensions

The suspensions filled in the microcups comprise a dielectric solvent with charged pigment particles dispersed therein and the particles migrate under the influence
10 of an electric field. The suspensions may optionally contain additional colorants which do not migrate in the electric field. The dispersion may be prepared according to methods well known in the art, such as US Patents Nos. 6,017,584, 5,914,806, 5,573,711, 5,403,518, 5,380,362, 4,680,103, 4,285,801, 4,093,534, 4,071,430, and 3,668,106, and as described in *IEEE Trans. Electron Devices*, ED-24, 827 (1977), and *J.*
15 *Appl. Phys.* 49(9), 4820 (1978).

The suspending fluid medium is a dielectric solvent which preferably has a low viscosity and a dielectric constant in the range of about 2 to about 30, preferably about 2 to about 15 for high particle mobility. Examples of suitable dielectric solvents include
20 hydrocarbons such as decahydronaphthalene (DECALIN), 5-ethylidene-2-norbornene, fatty oils, paraffin oil, aromatic hydrocarbons such as toluene, xylene, phenylxylylene, dodecylbenzene and alkyl naphthalene, halogenated solvents such as dichlorobenzotrifluoride, 3,4,5-trichlorobenzotrifluoride, chloropentafluoro-benzene, dichlorononane, pentachlorobenzene, and perfluoro solvents such as perfluorodecalin, perfluorotoluene, perfluoroxylene, FC-43, FC-70 and FC-5060 from 3M Company, St.
25 Paul MN, low molecular weight fluorine containing polymers such as poly(perfluoropropylene oxide) from TCI America, Portland, Oregon, poly(chlorotrifluoroethylene) such as Halocarbon Oils from Halocarbon Product Corp., River Edge, NJ, perfluoropolyalkylether such as Galden, HT-200, and Fluorolink from Ausimont or Krytox Oils and Greases K-Fluid Series from DuPont, Delaware. In one
30 preferred embodiment, poly(chlorotrifluoroethylene) is used as the dielectric solvent. In another preferred embodiment, poly(perfluoropropylene oxide) is used as the dielectric solvent.

The non-migrating fluid colorant may be formed from dyes or pigments. Nonionic azo and anthraquinone dyes are particularly useful. Examples of useful dyes include, but are not limited to: Oil Red EGN, Sudan Red, Sudan Blue, Oil Blue, Macrolex Blue, Solvent Blue 35, Pylam Spirit Black and Fast Spirit Black (Pylam Products Co., Arizona), Thermoplastic Black X-70 (BASF), anthraquinone blue, anthraquinone yellow 114, anthraquinone reds 111 and 135, anthraquinone green 28 and Sudan Black B (Aldrich). Fluorinated dyes are particularly useful when perfluorinated solvents are used. In the case of a pigment, the pigment particles for generating the non-migrating fluid colorant may also be dispersed in the dielectric solvent and these colored particles are preferably uncharged. If the pigment particles for generating the non-migrating fluid colorant are charged, they preferably carry a charge which is opposite from that of the charged migrating pigment particles. If both types of pigment particles carry the same charge, then they should have different charge density or different electrophoretic mobility. The dye or pigment for generating the non-migrating fluid colorant must be chemically stable and compatible with other components in the suspension.

The charged migrating pigment particles are preferably white, and may be organic or inorganic pigments, such as TiO_2 .

If colored migrating particles are used, they may be formed from phthalocyanine blue, phthalocyanine green, diarylide yellow, diarylide AAOT yellow, and quinacridone, azo, rhodamine, perylene pigment series (Sun Chemical), Hansa yellow G particles (Kanto Chemical), and Carbon Lampblack (Fisher). Submicron particle size is preferred. These particles should have acceptable optical characteristics, should not be swollen or softened by the dielectric solvent, and should be chemically stable. The resulting suspension must also be stable against sedimentation, creaming or flocculation under normal operating conditions.

The migrating pigment particles may exhibit a native charge, or may be charged explicitly using a charge control agent, or may acquire a charge when suspended in the dielectric solvent. Suitable charge control agents are well known in the art; they may be polymeric or non-polymeric in nature, and may also be ionic or non-ionic, including ionic surfactants such as Aerosol OT, sodium dodecylbenzenesulfonate, metal soaps, polybutene succinimide, maleic anhydride copolymers, vinylpyridine copolymers, vinylpyrrolidone copolymers (such as Ganex, International Specialty Products),

(meth)acrylic acid copolymers, and N,N-dimethylaminoethyl (meth)acrylate copolymers. Fluorosurfactants are particularly useful as charge controlling agents in perfluorocarbon solvents. These include FC fluorosurfactants such as FC-170C, FC-171, FC-176, FC430, FC431 and FC-740 from 3M Company and Zonyl
5 fluorosurfactants such as Zonyl FSA, FSE, FSN, FSN-100, FSO, FSO-100, FSD and UR from Dupont.

Suitable charged pigment dispersions may be manufactured by any of the well-known methods including grinding, milling, attriting, microfluidizing, and ultrasonic techniques. For example, pigment particles in the form of a fine powder are added to
10 the suspending solvent and the resulting mixture is ball milled or attrited for several hours to break up the highly agglomerated dry pigment powder into primary particles. Although less preferred, a dye or pigment for producing the non-migrating fluid colorant may be added to the suspension during the ball milling process.

Sedimentation or creaming of the pigment particles may be eliminated by
15 microencapsulating the particles with suitable polymers to match the specific gravity to that of the dielectric solvent. Microencapsulation of the pigment particles may be accomplished chemically or physically. Typical microencapsulation processes include interfacial polymerization, in-situ polymerization, phase separation, coacervation, electrostatic coating, spray drying, fluidized bed coating and solvent evaporation.

For pigment suspensions, there are many possibilities. For a subtractive color
20 system, the charged TiO₂ particles may be suspended in a dielectric fluid of cyan, yellow or magenta color. The cyan, yellow or magenta color may be generated via the use of a dye or a pigment. For an additive color system, the charged TiO₂ particles may be suspended in a dielectric solvent of red, green or blue color generated also via the use
25 of a dye or a pigment. The red, green, blue color system is preferred for most applications.

While the present invention has been described with reference to the specific embodiments thereof, it should be understood by those skilled in the art that various changes may be made and equivalents may be substituted without departing from the
30 true spirit and scope of the invention. In addition, many modifications may be made to adapt a particular situation, materials, compositions, processes, process step or steps, to the objective, spirit and scope of the present invention. All such modifications are intended to be within the scope of the claims appended hereto.

It is therefore wished that this invention to be defined by the scope of the appended claims as broadly as the prior art will permit, and in view of the specification.

WHAT IS CLAIMED IS:

1. An electrophoretic display comprising isolated electrophoretic cells formed from microcups of well-defined size, shape and aspect ratio with an in-plane switching mode.
- 5 2. The electrophoretic display of Claim 1 having two in-plane electrodes.
3. The electrophoretic display of Claim 1 having one in-plane electrode.
4. The electrophoretic display of Claim 1 wherein the microcups have a top opening ranging from about 10^2 to about $1 \times 10^6 \mu\text{m}^2$.
5. The electrophoretic display of Claim 4 wherein the microcups have a top
10 opening ranging from about 10^3 to about $1 \times 10^5 \mu\text{m}^2$.
6. The electrophoretic display of Claim 1 wherein the depth of the microcups is in the range of about 5 to about 200 microns.
7. The electrophoretic display of Claim 6 wherein the depth of the microcups is in the range of about 20 to about 100 microns.
- 15 8. The electrophoretic display of Claim 1 wherein the opening to the total area ratio is in the range of from about 0.2 to about 0.95.
9. An electrophoretic display having isolated cells formed from microcups of well-defined size, shape and aspect ratio, said cells comprising
a transparent viewing layer on one side of the display;
20 a layer having in-plane electrodes on the opposite side of the display; and
said cells are filled with a dielectric solvent or solvent mixture having charged particles dispersed therein.
10. The electrophoretic display of Claim 9 further comprising a separate background layer.
- 25 11. The electrophoretic display of Claim 9 wherein said transparent viewing layer is colorless.

12. The electrophoretic display of Claim 11 wherein said separate background layer is on top of the layer having in-plane electrodes.
13. The electrophoretic display of Claim 11 wherein said separate background layer is underneath the layer having in-plane electrodes.
- 5 14. The electrophoretic display of Claim 11 wherein said layer having in-plane electrodes serves as the background layer and the in-plane electrodes may be white or colored.
15. The electrophoretic display of Claim 9 wherein the display is a monochrome display.
- 10 16. The electrophoretic display of Claim 15 wherein said dielectric solvent is clear and colorless.
17. The electrophoretic display of Claim 16 wherein all said cells have particles of the white color and the same background color.
18. The electrophoretic display of Claim 17 wherein said background color is
15 black, red, green, blue, yellow, cyan or magenta.
19. The electrophoretic display of Claim 16 wherein all said cells have particles of the same color and the white background color.
20. The electrophoretic display of Claim 19 wherein said particles are black, red, green, blue, yellow, cyan or magenta.
- 20 21. The electrophoretic display of Claim 16 wherein said individual cells have particles of mixed colors and the same background color.
22. The electrophoretic display of Claim 21 wherein the mixed colors are two or more colors selected from a group consisting of black, white, red, green, blue, yellow, cyan and magenta.

23. The electrophoretic display of Claim 22 wherein said background color is selected from a group consisting of black, white, red, green, blue, yellow, cyan and magenta.

24. The electrophoretic display of Claim 9 wherein said display is a multiple
5 color display.

25. The electrophoretic display of Claim 24 wherein said cells have particles of the white color and different background colors.

26. The electrophoretic display of Claim 24 wherein said cells have particles of the black color and different background colors.

10 27. The electrophoretic display of Claim 24 wherein said cells have particles of different colors and a white background.

28. The electrophoretic display of Claim 24 wherein said cells have particles of different colors and a black background.

15 29. The electrophoretic display of Claim 9 wherein said transparent viewing layer is colored or by adding a color filter.

30. The electrophoretic display of Claim 29 wherein said separate background layer is on top of the layer having in-plane electrodes.

31. The electrophoretic display of Claim 29 wherein said separate background layer is underneath the layer having in-plane electrodes.

20 32. The electrophoretic display of Claim 29 wherein said layer having in-plane electrodes serves as the background layer and the in-plane electrodes may be white or colored.

33. The electrophoretic display of Claim 29 wherein all said cells have white particles and black background.

25 34. The electrophoretic display of Claim 33 wherein all said cells have the transparent viewing layer of the same color.

35. The electrophoretic display of Claim 33 wherein said cells have the transparent viewing layers of different colors.

36. A process for the manufacture of an electrophoretic display having isolated cells formed from microcups having well defined size, shape and aspect ratio,
5 said process comprises
- a) forming a layer by coating a radiation curable material over a transparent insulator substrate;
 - b) forming microcups on the radiation curable material by microembossing or imagewise exposure to radiation;
 - 10 c) filling the microcups with an electrophoretic composition;
 - d) sealing the microcups;
 - e) laminating the sealed microcups with a substrate containing in-plane electrodes.

37. The process for the manufacture of an electrophoretic display of Claim
15 36 with substrate containing in-plane electrodes and an adhesive.

38. A process for the manufacture of an electrophoretic display having isolated cells formed from microcups having well defined size, shape and aspect ratio, said process comprises
- a) forming a layer by coating a radiation curable material over a substrate
20 containing in-plane electrodes;
 - b) forming microcups on the radiation curable material by microembossing or imagewise exposure to radiation;
 - c) filling the microcups with an electrophoretic composition;
 - d) sealing the microcups;
 - 25 e) laminating the sealed microcups with a transparent insulator substrate.

39. The process for the manufacture of an electrophoretic display of Claim 38 with a transparent insulator substrate and an adhesive.

40. The process of Claim 36 wherein the microcups have a top opening ranging from about 10^2 to about $1 \times 10^6 \mu\text{m}^2$.

41. The process of Claim 38 wherein the microcups have a top opening ranging from about 10^2 to about $1 \times 10^6 \mu\text{m}^2$.

42. The process of Claim 36 wherein said transparent insulator substrate is colorless.

5 43. The process of Claim 42 wherein all said cells have the same background color and are filled with an electrophoretic composition comprising charged particles of the same color dispersed in a clear colorless dielectric solvent.

 44. The process of Claim 42 wherein said cells have different background colors and are filled with an electrophoretic composition comprising charged particles of
10 the same color dispersed in a clear colorless dielectric solvent.

 45. The process of Claim 42 wherein said cells have the same background color and are filled with an electrophoretic composition comprising charged particles of the different colors dispersed in a clear colorless dielectric solvent.

 46. The process of Claim 38 wherein said transparent insulator substrate is
15 colorless.

 47. The process of Claim 46 wherein all said cells have the same background color and are filled with an electrophoretic composition comprising charged particles of the same color dispersed in a clear colorless dielectric solvent.

 48. The process of Claim 46 wherein said cells have different background
20 colors and are filled with an electrophoretic composition comprising charged particles of the same color dispersed in a clear colorless dielectric solvent.

 49. The process of Claim 46 wherein said cells have the same background color and are filled with an electrophoretic composition comprising charged particles of different colors dispersed in a clear colorless dielectric solvent.

25 50. The process of Claim 36 wherein said transparent insulator substrate is colored.

51. The process of Claim 50 wherein said cells have the black background and are filled with an electrophoretic composition comprising charged white particles dispersed in a clear colorless dielectric solvent.

52. The process of Claim 51 wherein all cells have the transparent insulator
5 substrate of the same color.

53. The process of Claim 51 wherein said cells have the transparent insulator substrate of different colors.

54. The process of Claim 38 wherein said transparent insulator substrate is colored.

10 55. The process of Claim 54 wherein said cells have the black background and are filled with an electrophoretic composition comprising charged white particles dispersed in a clear colorless dielectric solvent.

56. The process of Claim 55 wherein all cells have the transparent insulator substrate of the same color.

15 57. The process of Claim 55 wherein said cells have the transparent insulator substrate of different colors.

58. The electrophoretic display of Claim 9 wherein a substrate containing an array of thin film transistors is used as the layer of in-plane electrodes.

1/3

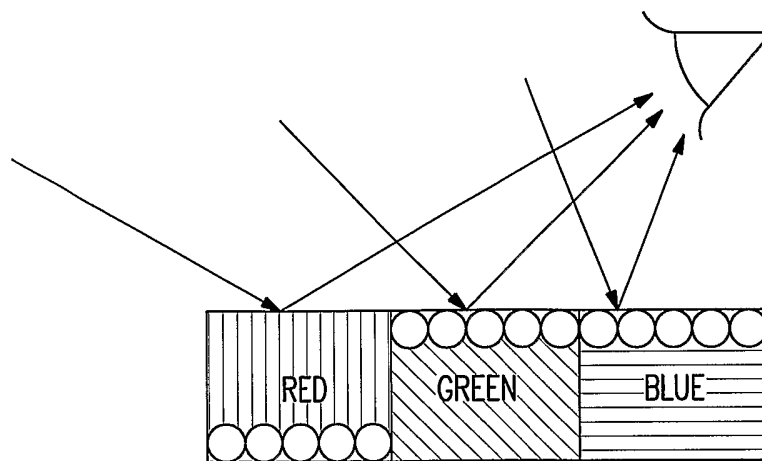


FIG. 1

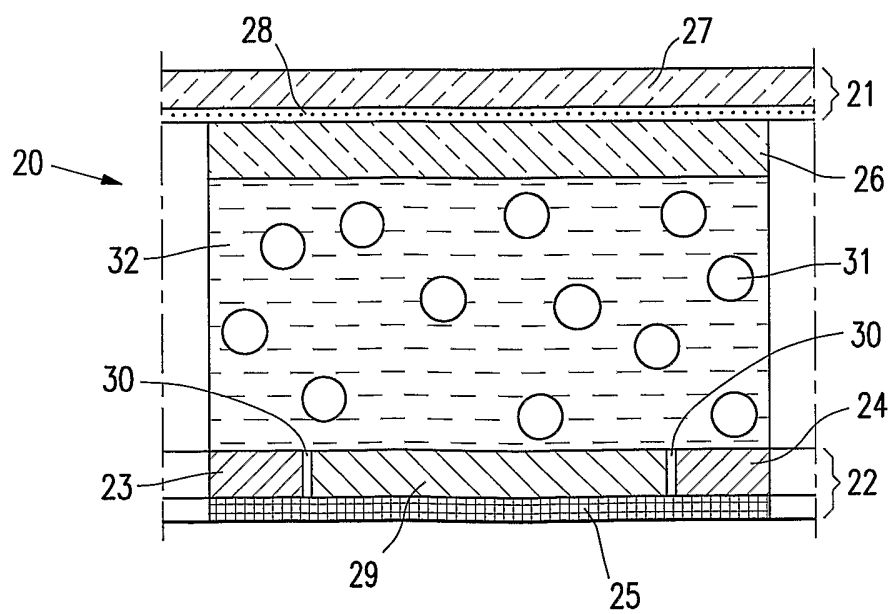
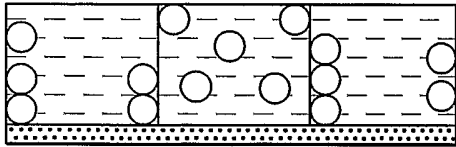
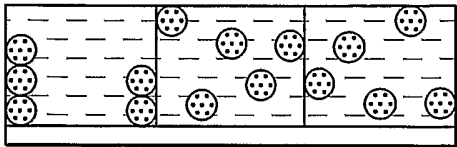


FIG. 2



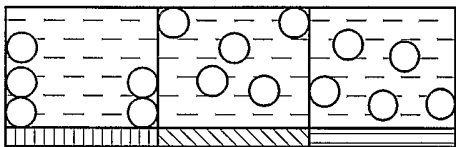
□ WHITE
◼ COLOR

FIG. 3A



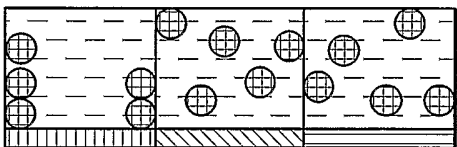
□ WHITE
◼ COLOR

FIG. 3B



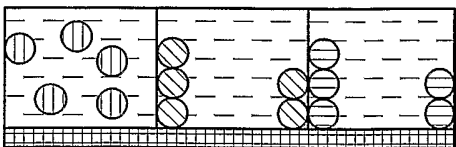
▤ RED
▥ GREEN
▧ BLUE
□ WHITE

FIG. 4A



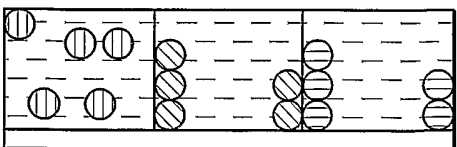
▤ RED
▥ GREEN
▧ BLUE
▩ BLACK

FIG. 4B



▤ RED
▥ GREEN
▧ BLUE
▩ BLACK

FIG. 4C



▤ RED
▥ GREEN
▧ BLUE
□ WHITE

FIG. 4D

3/3

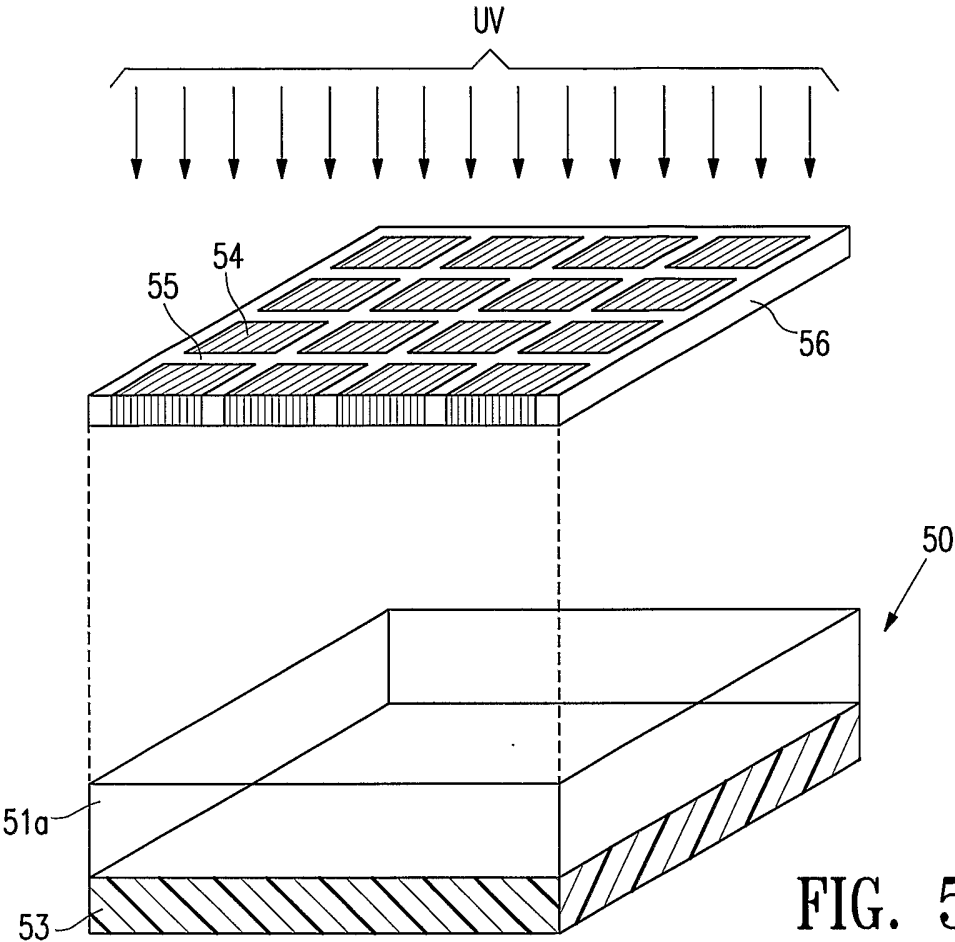


FIG. 5A

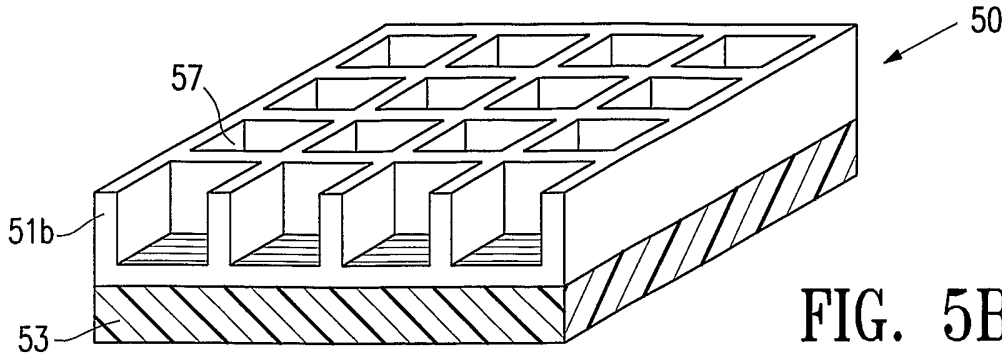


FIG. 5B

INTERNATIONAL SEARCH REPORT

Int ☐ national Application No

PCT/US 02/22812

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 G02F1/167

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

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, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	KISHI E ET AL: "DEVELOPMENT OF IN-PLANE EPD" 2000 SID INTERNATIONAL SYMPOSIUM DIGEST OF TECHNICAL PAPERS. LONG BEACH, CA, MAY 16 - 18, 2000, SID INTERNATIONAL SYMPOSIUM DIGEST OF TECHNICAL PAPERS, SAN JOSE, CA: SID, US, vol. 31, May 2000 (2000-05), pages 24-27, XP001086658 the whole document	1-58
A	HARBOUR ET AL: "Subdivided Electrophoretic Display" XEROX DISCLOSURE JOURNAL, XEROX CORPORATION. STAMFORD, CONN, US, vol. 4, no. 6, November 1979 (1979-11), page 705 XP002123212 the whole document	1-58
	--- -/--	

☒ 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.

& document member of the same patent family

Date of the actual completion of the international search

3 December 2002

Date of mailing of the international search report

09/12/2002

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

Gill, R

INTERNATIONAL SEARCH REPORT

Int  nal Application No

PCT/US 02/22812

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	PATENT ABSTRACTS OF JAPAN vol. 009, no. 026 (P-332), 5 February 1985 (1985-02-05) & JP 59 171930 A (MATSUSHITA DENKI SANGYO KK), 28 September 1984 (1984-09-28) abstract -----	1-58

INTERNATIONAL SEARCH REPORT

Information on patent family members

Internal Application No

PCT/US 02/22812

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
JP 59171930	A	28-09-1984	NONE