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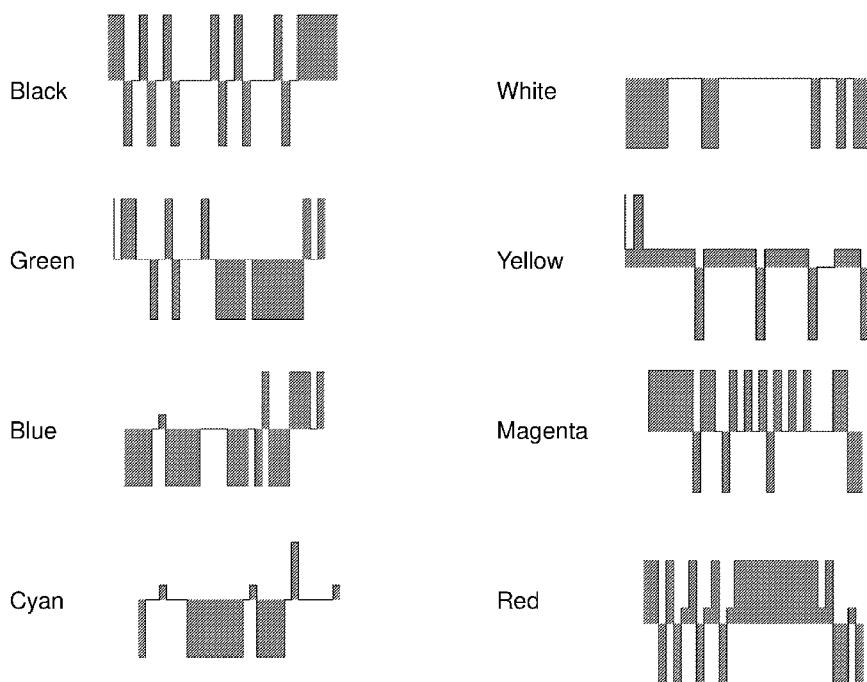


FIG. 6A

(57) Abstract: A color electrophoretic display includes a light-transmissive electrode at a viewing surface, a back electrode, and an electrophoretic medium disposed therebetween. The electrophoretic medium includes a non-polar fluid and a multi-pigment particle system having four types of charged electrophoretic pigment particles in the fluid. The particles includes first, second, third, and fourth types of particles having different optical properties. The second, third, and fourth types of particles have a charge polarity opposite to the charge polarity of the first type of particle. The multi-pigment particle system is directly addressable with push-pull waveforms applied to the back electrode having voltage levels selected from a set of exactly four different voltage levels to render any of eight primary colors (red, green, blue, cyan, magenta, yellow, black, and white) at each pixel while holding the voltage on the light-transmissive



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electrode constant. The voltage levels comprise a first positive voltage, a second lower positive voltage, a near zero voltage, and a negative voltage.

COLOR ELECTROPHORETIC MEDIUM HAVING FOUR PIGMENT PARTICLE SYSTEM ADDRESSABLE BY WAVEFORMS HAVING FOUR VOLTAGE LEVELS

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application claims priority from U.S. Provisional Patent Application No. 63,471,148 filed on June 5, 2023 entitled COLOR ELECTROPHORETIC MEDIUM HAVING FOUR PIGMENT PARTICLE SYSTEM ADDRESSABLE BY WAVEFORMS HAVING FOUR VOLTAGE LEVELS, which is hereby incorporated by reference herein in its entirety.

BACKGROUND

[0002] An electrophoretic display (EPD) changes color by modifying the position of a charged colored particle with respect to a light-transmissive viewing surface. Such electrophoretic displays are typically referred to as “electronic paper” or “ePaper” because the resulting display has high contrast and is sunlight-readable, much like ink on paper. Electrophoretic displays have enjoyed widespread adoption in eReaders, such as the AMAZON KINDLE® because the electrophoretic displays provide a book-like reading experience, use little power, and allow a user to carry a library of hundreds of books in a lightweight handheld device.

[0003] For many years, EPDs included only two types of charged color particles: black and white. (To be sure, “color” as used herein includes black and white.) The white particles are often of the light scattering type, and comprise, e.g., titanium dioxide, while the black particle are absorptive across the visible spectrum, and may comprise carbon black, or an absorptive metal oxide, such as copper chromite. In the simplest sense, a black and white EPD only requires a light-transmissive electrode at the viewing surface, a back electrode, and an electrophoretic medium including oppositely charged white and black particles. When a voltage of one polarity is provided, the white particles move to the viewing surface, and when a voltage of the opposite polarity is provided the black particles move to the viewing surface. If the back electrode includes controllable regions (pixels) – either segmented electrodes or an active matrix of pixel electrodes controlled by transistors – a pattern can be made to appear electronically at the viewing surface. The pattern can be, e.g., the text of a book.

[0004] More recently, a variety of color option have become commercially available for EPDs, including three-color displays (black, white, red; black white, yellow), four color displays (black, white, red, yellow), and color filter displays that rely on the black/white particles described above. EPDs with three or four reflective particles operate similar to the

conventional black and white displays because the desired color particle is driven to the viewing surface. The driving schemes are far more complicated than only black and white, but in the end, the optical function of the particles is the same, reflecting incoming light back to the viewer in the correct color.

[0005] Advanced Color electronic Paper (ACeP™) also included four particles, but the cyan, yellow, and magenta particles are subtractive rather than reflective, thereby allowing thousands of colors to be produced at each pixel. The color process is functionally equivalent to the printing methods that have long been used in offset printing and ink-jet printers. A given color is produced by using the correct ratio of cyan, yellow, and magenta on a bright white paper background. In the instance of ACeP, the relative positions of the cyan, yellow, magenta and white particles with respect to the viewing surface will determine the color at each pixel. While this type of EPD allows for thousands of colors at each pixel, it is critical to carefully control the position of each of the (50 to 500 nanometer-sized) pigments within a working space of about 10 to 20 micrometers in thickness. Obviously, variations in the position of the particles will result in incorrect colors being displayed at a given pixel. Accordingly, exquisite voltage control is required for such a system. More details of this system are available in the following U.S. patents, all of which are incorporated by reference in their entireties: U.S. Patent Nos. 9,361,836; 9,921,451; 10,276,109; 10,353,266; 10,467,984; and 10,593,272.

[0006] The term gray state is used herein in its conventional meaning in the imaging art to refer to a state intermediate two extreme optical states of a pixel, and does not necessarily imply a black-white transition between these two extreme states. For example, several of the E Ink patents and published applications referred to below describe EPDs in which the extreme states are white and deep blue, so that an intermediate gray state would actually be pale blue. Indeed, as already mentioned, the change in optical state may not be a color change at all. The terms black and white may be used hereinafter to refer to the two extreme optical states of a display, and should be understood as normally including extreme optical states that are not strictly black and white, e.g., the aforementioned white and dark blue states.

[0007] The terms bistable and bistability are used herein in their conventional meaning in the art to refer to displays comprising display elements having first and second display states differing in at least one optical property, and such that after any given element has been driven, by means of an addressing pulse of finite duration, to assume either its first or second display state, after the addressing pulse has terminated, that state will persist for at least several times, e.g., at least four times, the minimum duration of the addressing pulse required to change the

state of the display element. It is shown in U.S. Patent No. 7,170,670 that some particle-based EPDs capable of gray scale are stable not only in their extreme black and white states but also in their intermediate gray states, and the same is true of some other types of electro-optic displays. This type of display is properly called multi-stable rather than bistable, although for convenience the term bistable may be used herein to cover both bistable and multi-stable displays.

[0008] The term impulse, when used to refer to driving an EPD, is used herein to refer to the integral of the applied voltage with respect to time during the period in which the display is driven.

[0009] A particle that absorbs, scatters, or reflects light, either in a broad band or at selected wavelengths, is referred to herein as a colored or pigment particle. Various materials other than pigments (in the strict sense of that term as meaning insoluble colored materials) that absorb or reflect light, such as dyes or photonic crystals, etc., may also be used in the electrophoretic media and displays of the present invention.

[0010] Particle-based EPDs have been the subject of intense research and development for a number of years. In such displays, a plurality of charged particles (sometimes referred to as pigment particles) move through a fluid under the influence of an electric field. EPDs can have attributes of good brightness and contrast, wide viewing angles, state bistability, and low power consumption when compared with liquid crystal displays. Nevertheless, problems with the long-term image quality of these displays have prevented their widespread usage. For example, particles that make up EPDs tend to settle, resulting in inadequate service-life for these displays.

[0011] As noted above, electrophoretic media require the presence of a fluid. In most prior art electrophoretic media, this fluid is a liquid, but electrophoretic media can be produced using gaseous fluids; see, e.g., Kitamura, T., et al., Electrical toner movement for electronic paper-like display, IDW Japan, 2001, Paper HCS1-1, and Yamaguchi, Y., et al., Toner display using insulative particles charged triboelectrically, IDW Japan, 2001, Paper AMD4-4). See also U.S. Patents Nos. 7,321,459 and 7,236,291. Such gas-based electrophoretic media appear to be susceptible to the same types of problems due to particle settling as liquid-based electrophoretic media, when the media are used in an orientation which permits such settling, e.g., in a sign where the medium is disposed in a vertical plane. Indeed, particle settling appears to be a more serious problem in gas-based electrophoretic media than in liquid-based ones, since the lower viscosity of gaseous suspending fluids as compared with liquid ones allows more rapid settling

of the electrophoretic particles.

[0012] Numerous patents and applications assigned to or in the names of the Massachusetts Institute of Technology (MIT) and E Ink Corporation describe various technologies used in encapsulated electrophoretic and other electro-optic media. Such encapsulated media comprise numerous small capsules, each of which itself comprises an internal phase containing electrophoretically-mobile particles in a fluid medium, and a capsule wall surrounding the internal phase. Typically, the capsules are themselves held within a polymeric binder to form a coherent layer positioned between two electrodes. The technologies described in these patents and applications include:

- (a) Electrophoretic particles, fluids and fluid additives; see, e.g., U.S. Patent Nos. 7,002,728 and 7,679,814;
- (b) Capsules, binders and encapsulation processes; see, e.g., U.S. Patents Nos. 6,922,276 and 7,411,719;
- (c) Microcell structures, wall materials, and methods of forming microcells; see, e.g., United States Patent Nos. 7,072,095 and 9,279,906;
- (d) Methods for filling and sealing microcells; see, e.g., United States Patent Nos. 7,144,942 and 7,715,088;
- (e) Films and sub-assemblies containing electro-optic materials; see, e.g., U.S. Patents Nos. 6,982,178 and 7,839,564;
- (f) Backplanes, adhesive layers and other auxiliary layers and methods used in displays; see, e.g., U.S. Patents Nos. 7,116,318 and 7,535,624;
- (g) Color formation color adjustment; see, e.g., U.S. Patent Nos. 6,017,584; 6,545,797; 6,664,944; 6,788,452; 6,864,875; 6,914,714; 6,972,893; 7,038,656; 7,038,670; 7,046,228; 7,052,571; 7,075,502***; 7,167,155; 7,385,751; 7,492,505; 7,667,684; 7,684,108; 7,791,789; 7,800,813; 7,821,702; 7,839,564***; 7,910,175; 7,952,790; 7,956,841; 7,982,941; 8,040,594; 8,054,526; 8,098,418; 8,159,636; 8,213,076; 8,363,299; 8,422,116; 8,441,714; 8,441,716; 8,466,852; 8,503,063; 8,576,470; 8,576,475; 8,593,721; 8,605,354; 8,649,084; 8,670,174; 8,704,756; 8,717,664; 8,786,935; 8,797,634; 8,810,899; 8,830,559; 8,873,129; 8,902,153; 8,902,491; 8,917,439; 8,964,282; 9,013,783; 9,116,412; 9,146,439; 9,164,207; 9,170,467; 9,170,468; 9,182,646; 9,195,111; 9,199,441; 9,268,191; 9,285,649; 9,293,511; 9,341,916; 9,360,733; 9,361,836; 9,383,623; and 9,423,666; and U.S. Patent Applications Publication Nos.

2008/0043318; 2008/0048970; 2009/0225398; 2010/0156780; 2011/0043543; 2012/0326957; 2013/0242378; 2013/0278995; 2014/0055840; 2014/0078576; 2014/0340430; 2014/0340736; 2014/0362213; 2015/0103394; 2015/0118390; 2015/0124345; 2015/0198858; 2015/0234250; 2015/0268531; 2015/0301246; 2016/0011484; 2016/0026062; 2016/0048054; 2016/0116816; 2016/0116818; and 2016/0140909;

(h) Methods for driving displays; see, e.g., U.S. Patent Nos. 5,930,026; 6,445,489; 6,504,524; 6,512,354; 6,531,997; 6,753,999; 6,825,970; 6,900,851; 6,995,550; 7,012,600; 7,023,420; 7,034,783; 7,061,166; 7,061,662; 7,116,466; 7,119,772; 7,177,066; 7,193,625; 7,202,847; 7,242,514; 7,259,744; 7,304,787; 7,312,794; 7,327,511; 7,408,699; 7,453,445; 7,492,339; 7,528,822; 7,545,358; 7,583,251; 7,602,374; 7,612,760; 7,679,599; 7,679,813; 7,683,606; 7,688,297; 7,729,039; 7,733,311; 7,733,335; 7,787,169; 7,859,742; 7,952,557; 7,956,841; 7,982,479; 7,999,787; 8,077,141; 8,125,501; 8,139,050; 8,174,490; 8,243,013; 8,274,472; 8,289,250; 8,300,006; 8,305,341; 8,314,784; 8,373,649; 8,384,658; 8,456,414; 8,462,102; 8,514,168; 8,537,105; 8,558,783; 8,558,785; 8,558,786; 8,558,855; 8,576,164; 8,576,259; 8,593,396; 8,605,032; 8,643,595; 8,665,206; 8,681,191; 8,730,153; 8,810,525; 8,928,562; 8,928,641; 8,976,444; 9,013,394; 9,019,197; 9,019,198; 9,019,318; 9,082,352; 9,171,508; 9,218,773; 9,224,338; 9,224,342; 9,224,344; 9,230,492; 9,251,736; 9,262,973; 9,269,311; 9,299,294; 9,373,289; 9,390,066; 9,390,661; and 9,412,314; and U.S. Patent Applications Publication Nos. 2003/0102858; 2004/0246562; 2005/0253777; 2007/0091418; 2007/0103427; 2007/0176912; 2008/0024429; 2008/0024482; 2008/0136774; 2008/0291129; 2008/0303780; 2009/0174651; 2009/0195568; 2009/0322721; 2010/0194733; 2010/0194789; 2010/0220121; 2010/0265561; 2010/0283804; 2011/0063314; 2011/0175875; 2011/0193840; 2011/0193841; 2011/0199671; 2011/0221740; 2012/0001957; 2012/0098740; 2013/0063333; 2013/0194250; 2013/0249782; 2013/0321278; 2014/0009817; 2014/0085355; 2014/0204012; 2014/0218277; 2014/0240210; 2014/0240373; 2014/0253425; 2014/0292830; 2014/0293398; 2014/0333685; 2014/0340734; 2015/0070744; 2015/0097877; 2015/0109283; 2015/0213749; 2015/0213765; 2015/0221257; 2015/0262255; 2015/0262551; 2016/0071465; 2016/0078820; 2016/0093253; 2016/0140910; and 2016/0180777 (these patents and applications may

hereinafter be referred to as the MEDEOD (MEthods for Driving Electro-optic Displays) applications);

(i) Applications of displays; see, e.g., U.S. Patents Nos. 7,312,784 and 8,009,348; and

(j) Non-electrophoretic displays, as described in U.S. Patents Nos. 6,241,921; and U.S. Patent Applications Publication Nos. 2015/0277160; and U.S. Patent Application Publications Nos. 2015/0005720 and 2016/0012710.

[0013] Many of the aforementioned patents and applications recognize that the walls surrounding the discrete microcapsules in an encapsulated electrophoretic medium could be replaced by a continuous phase, thus producing a so-called polymer-dispersed EPD, in which the electrophoretic medium comprises a plurality of discrete droplets of an electrophoretic fluid and a continuous phase of a polymeric material, and that the discrete droplets of electrophoretic fluid within such a polymer-dispersed EPD may be regarded as capsules or microcapsules even though no discrete capsule membrane is associated with each individual droplet; see, e.g., U.S. Patent No. 6,866,760. Accordingly, for purposes of the present application, such polymer-dispersed electrophoretic media are regarded as sub-species of encapsulated electrophoretic media.

[0014] A related type of EPD is a so-called microcell EPD. In a microcell EPD, the charged particles and the fluid are not encapsulated within microcapsules, but instead are retained within a plurality of cavities formed within a carrier medium, typically a polymeric film. See, e.g., U.S. Patents Nos. 6,672,921 and 6,788,449.

[0015] Although electrophoretic media are often opaque (since, e.g., in many electrophoretic media, the particles substantially block transmission of visible light through the display) and operate in a reflective mode, many EPDs can be made to operate in a so-called shutter mode in which one display state is substantially opaque and one is light-transmissive. See, e.g., U.S. Patents Nos. 5,872,552; 6,130,774; 6,144,361; 6,172,798; 6,271,823; 6,225,971; and 6,184,856. Dielectrophoretic displays, which are similar to EPDs but rely upon variations in electric field strength, can operate in a similar mode. See U.S. Patent No. 4,418,346. Other types of electro-optic displays may also be capable of operating in shutter mode. Electro-optic media operating in shutter mode can be used in multi-layer structures for full color displays; in such structures, at least one layer adjacent the viewing surface of the display operates in shutter mode to expose or conceal a second layer more distant from the viewing surface.

[0016] An encapsulated EPD typically does not suffer from the clustering and settling

failure mode of traditional electrophoretic devices and provides further advantages, such as the ability to print or coat the display on a wide variety of flexible and rigid substrates. (Use of the word printing is intended to include all forms of printing and coating, including, but without limitation: pre-metered coatings such as patch die coating, slot or extrusion coating, slide or cascade coating, curtain coating; roll coating such as knife over roll coating, forward and reverse roll coating; gravure coating; dip coating; spray coating; meniscus coating; spin coating; brush coating; air knife coating; silk screen printing processes; electrostatic printing processes; thermal printing processes; ink jet printing processes; electrophoretic deposition (See U.S. Patent No. 7,339,715); and other similar techniques.) Thus, the resulting display can be flexible. Further, because the display medium can be printed (using a variety of methods), the display itself can be made inexpensively.

[0017] As indicated above, most simple prior art electrophoretic media essentially display only two colors. Such electrophoretic media either use a single type of electrophoretic particle having a first color in a colored fluid having a second, different color (in which case, the first color is displayed when the particles lie adjacent the viewing surface of the display and the second color is displayed when the particles are spaced from the viewing surface), or first and second types of electrophoretic particles having differing first and second colors in an uncolored fluid (in which case, the first color is displayed when the first type of particles lie adjacent the viewing surface of the display and the second color is displayed when the second type of particles lie adjacent the viewing surface). Typically the two colors are black and white. If a full color display is desired, a color filter array may be deposited over the viewing surface of the monochrome (black and white) display. Displays with color filter arrays rely on area sharing and color blending to create color stimuli. The available display area is shared between three or four primary colors such as red/green/blue (RGB) or red/green/blue/white (RGBW), and the filters can be arranged in one-dimensional (stripe) or two-dimensional (2x2) repeat patterns. Other choices of primary colors or more than three primaries are also known in the art. The three (in the case of RGB displays) or four (in the case of RGBW displays) sub-pixels are chosen small enough so that at the intended viewing distance they visually blend together to a single pixel with a uniform color stimulus ('color blending'). The inherent disadvantage of area sharing is that the colorants are always present, and colors can only be modulated by switching the corresponding pixels of the underlying monochrome display to white or black (switching the corresponding primary colors on or off). For example, in an ideal RGBW display, each of the red, green, blue and white primaries occupy one fourth of the display area

(one sub-pixel out of four), with the white sub-pixel being as bright as the underlying monochrome display white, and each of the colored sub-pixels being no lighter than one third of the monochrome display white. The brightness of the white color shown by the display as a whole cannot be more than one half of the brightness of the white sub-pixel (white areas of the display are produced by displaying the one white sub-pixel out of each four, plus each colored sub-pixel in its colored form being equivalent to one third of a white sub-pixel, so the three colored sub-pixels combined contribute no more than the one white sub-pixel). The brightness and saturation of colors is lowered by area-sharing with color pixels switched to black. Area sharing is especially problematic when mixing yellow because it is lighter than any other color of equal brightness, and saturated yellow is almost as bright as white. Switching the blue pixels (one fourth of the display area) to black makes the yellow too dark.

[0018] U.S. Patent Nos. 8,576,476 and 8,797,634 describe multicolor EPDs having a single back plane comprising independently addressable pixel electrodes and a common, light-transmissive front electrode. A plurality of electrophoretic layers are disposed between the back plane and the front electrode. Displays described in these patents are capable of rendering any of the primary colors (red, green, blue, cyan, magenta, yellow, white and black) at any pixel location. However, there are disadvantages to the use of multiple electrophoretic layers located between a single set of addressing electrodes. The electric field experienced by the particles in a particular layer is lower than would be the case for a single electrophoretic layer addressed with the same voltage. In addition, optical losses in an electrophoretic layer closest to the viewing surface (e.g., caused by light scattering or unwanted absorption) may affect the appearance of images formed in underlying electrophoretic layers.

[0019] Two other types of EPD systems provide a single electrophoretic medium capable of rendering any color at any pixel location. Specifically, U.S. Patent No. 9,697,778 describes a display in which a dyed solvent is combined with a white (light-scattering) particle that moves in a first direction when addressed with a low applied voltage and in the opposite direction when addressed with a higher voltage. When the white particles and the dyed solvent are combined with two additional particles of opposite charge to the white particle, it is possible to render a full-color display. However, the color states of the '778 patent are not acceptable for applications such as a text reader. In particular, there will always be some of the dyed fluid separating the white scattering particle from the viewing surface, which leads to a tint in the white state of the display.

[0020] A second form of electrophoretic medium capable of rendering any color at any

pixel location is described in U.S. Patent No. 9,921,451. In the '451 patent, the electrophoretic medium includes four particles: white, cyan, magenta and yellow, in which two of the particles are positively-charged and two negatively charged. However displays of the '451 patent also suffer from color mixing with the white state. Because one of the particles has the same charge as the white particle, some quantity of the same-charge particle moves with the white toward the viewing surface when the white state is desired. While it is possible to overcome this unwanted tinting with complex waveforms driving the display, such waveforms greatly increase the update time of the display and in some instances, result in unacceptable "flashing" between images.

SUMMARY

[0021] Disclosed herein are improved full-color EPDs that are addressable by driving waveforms having four voltage levels. The EPDs include an electrophoretic medium having four types of charged electrophoretic pigment particles including three substantially non-light-scattering pigments that are colored, preferably corresponding to the subtractive primary colors (cyan, magenta, and yellow), and a fourth, light-scattering white pigment. The three colored pigments have one charge polarity, while the white pigment has the opposite charge polarity. Preferably the white pigment is negatively charged, and the three colored pigments are positively charged. The display is addressed by waveforms having exactly four different voltage levels to render any of the primary colors (white, black, cyan, magenta, yellow, red, green, and blue) at any pixel location. The voltage levels preferably comprise high and low positive voltages, a single negative voltage, and a near zero voltage. Because the EPD utilizes exactly four voltage levels and can be addressed by two bits of data per pixel (instead of three or more bits used in prior art devices), device costs can be reduced while providing excellent color quality without significant sacrifice of color gamut.

[0022] In a first aspect, the invention provides a color electrophoretic display, comprising a light-transmissive electrode at a viewing surface, a back electrode, and an electrophoretic medium disposed between the light-transmissive electrode and the back electrode. The electrophoretic medium comprises a non-polar fluid and a multi-pigment particle system comprising four types of charged electrophoretic pigment particles dispersed in the non-polar fluid. The four types of charged electrophoretic pigment particles comprise a first type of particle having a first optical property and a first charge polarity; a second type of particle having a second optical property and a second charge polarity opposite to the first charge

polarity; a third type of particle having a third optical property and the second charge polarity; and a fourth type of particle having a fourth optical property and the second charge polarity. The first, second, third, and fourth optical properties are different from one another. The multi-pigment particle system is directly addressable with push-pull waveforms applied to the back electrode having pulse voltage levels selected from a set of exactly four different voltage levels to render any of eight primary colors of red, green, blue, cyan, magenta, yellow, black, and white at each pixel of the electrophoretic medium while holding the voltage on the light-transmissive electrode constant. The voltage levels comprise a first positive voltage, a lower second positive voltage, a near zero voltage, and a negative voltage.

[0023] In a second aspect, the invention provides a method, comprising providing a color electrophoretic display, comprising a light-transmissive electrode at a viewing surface, a back electrode, and an electrophoretic medium disposed between the light-transmissive electrode and the back electrode. The electrophoretic medium comprises a non-polar fluid and a multi-pigment particle system comprising four types of charged electrophoretic pigment particles dispersed in the non-polar fluid. The four types of charged electrophoretic pigment particles comprise: a first type of particle having a first optical property and a first charge polarity; a second type of particle having a second optical property and a second charge polarity, wherein the second charge polarity is opposite to the first charge polarity; a third type of particle having a third optical property and the second charge polarity; and a fourth type of particle having a fourth optical property and the second charge polarity. The first, second, third, and fourth optical properties are different from one another. The method also includes directly addressing the multi-pigment particle system with push-pull waveforms applied to the back electrode having voltage levels selected from a set of exactly four different voltage levels to render any of eight primary colors of red, green, blue, cyan, magenta, yellow, black, and white at each pixel of the electrophoretic medium while holding the voltage on the light-transmissive electrode constant. The four different voltage levels comprise a first positive voltage, a lower second positive voltage, a near zero voltage, and a negative voltage.

[0024] In one or more embodiments, the first positive voltage is 15 to 30V, the second positive voltage is 5 to 15V, and the negative voltage is -15 to -30V.

[0025] In one or more embodiments, the first positive voltage is 24V, the second positive voltage is 10V, and the negative voltage is -24V.

[0026] In one or more embodiments, the push-pull waveforms comprise aperiodic voltage

sequences.

[0027] In one or more embodiments, the first type of particle is a light scattering particle, and the second, third, and fourth types of particles are light-absorbing particles.

[0028] In one or more embodiments, the first type of particle is white, and the second, third, and fourth types of particles are selected from cyan, magenta, and yellow.

[0029] In one or more embodiments, the first type of particle is white, the second type of particle is cyan, the third type of particle is magenta, and the fourth type of particle is yellow.

[0030] In one or more embodiments, the first charge polarity is negative, and the second charge polarity is positive.

[0031] In one or more embodiments, the multi-pigment particle system has exactly four types of charged electrophoretic particles.

[0032] In one or more embodiments, the non-polar fluid includes charge-control additives.

[0033] In one or more embodiments, the electrophoretic medium is encapsulated in capsules or contained in sealed microcells.

[0034] In one or more embodiments, the electrophoretic display is configured for incorporation in a book reader, a portable computer, a tablet computer, a monitor, a phone, a smart card, a sign, a watch, jewelry, a shelf label, a panel for a vehicle, or a flash drive.

[0035] In one or more embodiments, the back electrode comprises a segmented electrode or a backplane including an array of pixel electrodes.

[0036] In one or more embodiments, the backplane includes an array of thin film transistors coupled to the pixel electrodes.

[0037] In one or more embodiments, the backplane includes an array of thin film transistors coupled to the pixel electrodes.

[0038] In one or more embodiments, the push-pull waveforms are determined using a trained computer model.

[0039] In one or more embodiments, a system is disclosed comprising a color electrophoretic display in which the back electrode comprises a backplane including an array of pixel electrodes. The system further includes a source driver; at least one power supply; and a controller for transmitting two bits of data per pixel to the source driver for controlling the source driver to apply selected voltages from the at least one power supply to selected pixel electrodes.

BRIEF DESCRIPTION OF DRAWINGS

[0040] FIG. 1 is a schematic cross-section showing the positions of the various colored particles in an electrophoretic medium when displaying black, white, the three subtractive primary colors, and the three additive primary colors.

[0041] FIG. 2A is a simplified illustration of an exemplary EPD having four types of particles (white, yellow, magenta, and cyan) in a non-polar fluid capable of rendering a full range of colors at each pixel electrode.

[0042] FIG. 2B illustrates a transition between a first optical state having all of the particles of a first charge polarity at the viewing surface and a second optical state having the particles with the second (opposite) polarity at the viewing surface.

[0043] FIG. 2C illustrates a transition between a first optical state having all of the particles of the first charge polarity at the viewing surface and a third optical state having the particles with the second (opposite) polarity behind the middle charged particles of the first polarity, which are located at the viewing surface.

[0044] FIG. 2D illustrates a transition between a first optical state having all of the particles of the first charge polarity at the viewing surface and a fourth optical state having the particles with the second (opposite) polarity behind the low charged particles of the first polarity, which are located at the viewing surface.

[0045] FIG. 2E illustrates a transition between a first optical state having all of the particles of the first charge polarity at the viewing surface and a fifth optical state having the particles with the second (opposite) polarity behind a combination of the low charged particles and the medium charged particles of the first polarity, which are located at the viewing surface.

[0046] FIG. 3 illustrates an exemplary equivalent circuit of a single pixel of an EPD.

[0047] FIG. 4 is a simplified illustration showing the layers of an exemplary EPD.

[0048] FIG. 5 shows exemplary push-pull drive waveforms having five voltage levels for addressing a four-particle electrophoretic medium having white, yellow, magenta, and cyan particles.

[0049] FIGS. 6A and 6B show exemplary push-pull drive waveforms having four voltage levels for addressing a four-particle color electrophoretic medium in accordance with one or more embodiments.

[0050] FIG. 7 is a diagrammatic view of an exemplary driving system for controlling a four-particle color electrophoretic medium in accordance with one or more embodiments.

[0051] FIG. 8 is a graph comparing the color gamut volume resulting from waveforms

having different numbers of driving voltage levels.

[0052] FIG. 9 shows the color gamut available using waveforms having different numbers of driving voltage levels.

DETAILED DESCRIPTION

[0053] Various embodiments of the invention disclosed herein relate to a full-color EPD that is addressable by driving waveforms having four voltage levels. The EPD includes an electrophoretic medium having four types of charged electrophoretic pigment particles including three substantially non-light-scattering pigments that are colored, preferably corresponding to the subtractive primary colors (cyan, magenta, and yellow), and a fourth, light-scattering white pigment. The three colored pigments have one charge polarity, while the white pigment has the opposite charge polarity. Preferably the white pigment is negatively charged, and the three colored pigments are positively charged. The display is addressed by waveforms having exactly four different voltage levels to render any of the primary colors (white, black, cyan, magenta, yellow, red, green, and blue) at any pixel location. The voltage levels are specified by two bits of data and preferably comprise high and low positive voltages, a single negative voltage, and a near zero voltage. As used herein, a “near zero voltage” means -0.5 V to 0.5 V. Because the EPD utilizes exactly four voltage levels and can be addressed by 2 bits of data per pixel, instead of 3 or more bits used in prior art devices, device costs can be reduced while providing excellent color quality without significant sacrifice of color gamut, as discussed below.

[0054] By way of background, U.S. Patent Application Publication No. 20220082896, the content of which is incorporated herein by reference in its entirety, discloses a four-particle electrophoretic medium, including a first particle of a first polarity and three other particles having the opposite polarity and different magnitudes of charge. Typically, such a system includes a negative white particle and yellow, magenta, and cyan positively-charged particles having subtractive primary colors. Additionally, some particles may be engineered so that their electrophoretic mobility is non-linear with respect to the strength of the applied electric field. Accordingly, one or more particles will experience a decrease in electrophoretic mobility with the application of a high electric field (e.g., 20V or greater) of the correct polarity. Such a four-particle system is shown schematically in FIG. 1, and it can provide white, yellow, red,

magenta, blue, cyan, green, and black at every pixel.

[0055] As shown in FIG. 1, each of the eight principal colors (red, green, blue, cyan, magenta, yellow, black, and white) corresponds to a different arrangement of the four particles, such that the viewer only sees those colored particles that are on the viewing side of the white particle (i.e., the only particle that scatters light). To achieve a wide range of colors, additional voltage levels are used for finer control of the particles. In the formulations described, the first (typically negative) particle is reflective (typically white), while the other three particles oppositely charged (typically positive) particles include three substantially non-light-scattering (“SNLS”). The use of SNLS particles allows mixing of colors and provides for more color outcomes than can be achieved with the same number of scattering particles. These thresholds must be sufficiently separated for avoidance of cross-talk, and this separation necessitates the use of high addressing voltages for some colors. The four-particle electrophoretic media can also be updated faster, require “less flashy” transitions, and produce color spectra that is more pleasing to the viewer (and thus, commercially more valuable). Additionally, the disclosed formulations provides for fast (e.g., less than 500 ms, e.g., less than 300 ms, e.g., less than 200 ms, e.g., less than 100 ms) updates between black and white pixels, thereby enabling fast page turns for black on white text.

[0056] In FIG. 1, it is assumed that the viewing surface of the display is at the top (as illustrated), i.e., a user views the display from this direction, and light is incident from this direction. As already noted, only one of the four particles used in the electrophoretic medium substantially scatters light, and in FIG. 1 this particle is assumed to be the white pigment. This light-scattering white particle forms a white reflector against which any particles above the white particles (as illustrated in FIG. 1) are viewed. Light entering the viewing surface of the display passes through these particles, is reflected from the white particles, passes back through these particles and emerges from the display. Thus, the particles above the white particles may absorb various colors and the color appearing to the user is that resulting from the combination of particles above the white particles. Any particles disposed below (behind from the user’s point of view) the white particles are masked by the white particles and do not affect the color displayed. Because the second, third and fourth particles are substantially non-light-scattering, their order or arrangement relative to each other is unimportant, but for reasons already stated, their order or arrangement with respect to the white (light-scattering) particles is critical.

[0057] More specifically, when the cyan, magenta, and yellow particles lie below the white particles (Situation [A] in FIG. 1), there are no particles above the white particles and the pixel

simply displays a white color. When a single particle is above the white particles, the color of that single particle is displayed, yellow, magenta and cyan in Situations [B], [D], and [F], respectively in FIG. 1. When two particles lie above the white particles, the color displayed is a combination of those of these two particles; in FIG. 1, in Situation [C], magenta and yellow particles display a red color, in Situation [E], cyan and magenta particles display a blue color, and in Situation [G], yellow and cyan particles display a green color. Finally, when all three colored particles lie above the white particles (Situation [H] in FIG. 1), all the incoming light is absorbed by the three subtractive primary colored particles and the pixel displays a black color.

[0058] It is possible that one subtractive primary color could be rendered by a particle that scatters light, so that the display would comprise two types of light-scattering particles, one of which would be white and the other colored. In this case, however, the position of the light-scattering colored particle with respect to the other colored particles overlying the white particle would be important. For example, in rendering the color black (when all three colored particles lie over the white particles), the scattering colored particle cannot lie over the non-scattering colored particles (otherwise they will be partially or completely hidden behind the scattering particle and the color rendered will be that of the scattering colored particle, not black).

[0059] FIG. 1 shows an idealized situation in which the colors are uncontaminated (i.e., the light-scattering white particles completely mask any particles lying behind the white particles). In practice, the masking by the white particles may be imperfect so that there may be some small absorption of light by a particle that ideally would be completely masked. Such contamination typically reduces both the lightness and the chroma of the color being rendered. Such color contamination should be minimized to the point that the colors formed are commensurate with an industry standard for color rendition. A particularly favored standard is SNAP (the standard for newspaper advertising production), which specifies L^* , a^* , and b^* values for each of the eight primary colors referred to above. (Hereinafter, “primary colors” will be used to refer to the eight colors: black, white, the three subtractive primaries, and the three additive primaries as shown in FIG. 1.)

[0060] FIGS. 2A-2E, which are also disclosed in U.S. Patent Application Publication No. 20220082896, show schematic cross-sectional representations of the four particle types. A display layer utilizing the improved electrophoretic medium includes a first (viewing) surface 13 on the viewing side, and a second surface 14 on the opposite side of the first surface 13. The

electrophoretic medium is disposed between the two surfaces. Each space between two dotted vertical lines denotes a pixel. Within each pixel, the electrophoretic medium can be addressed and the viewing surface 13 of each pixel can achieve the color states shown in FIG. 1 without a need for additional layers, and without a color filter array.

[0061] As standard with EPDs, the first surface 13 includes a common electrode 11, which is light-transmissive, e.g., constructed from a sheet of PET with indium tin oxide (ITO) disposed thereon. On the second surface 14, there is an electrode layer 12, which includes a plurality of pixel electrodes 15. Such pixel electrodes are described in U.S. Patent No. 7,046,228, the content of which is incorporated herein by reference in its entirety. It is noted that while active matrix driving with a thin film transistor (TFT) backplane is mentioned for the layer of pixel electrodes, other types of electrode addressing can also be used as long as the electrodes serve the desired functions. For example, the top and bottom electrodes can be contiguous or segmented. Additionally, pixel electrode backplanes different from those described in the '228 patent are also suitable, and may include active matrix backplanes capable of providing higher driving voltages than typically found with amorphous silicon thin-film-transistor backplanes.

[0062] Newly-developed active matrix backplanes include thin film transistors incorporating metal oxide materials, such as tungsten oxide, tin oxide, indium oxide, zinc oxide or more complex metal oxides such as indium gallium zirconium oxide. In these applications, a channel formation region is formed for each transistor using such metal oxide materials, allowing faster switching of higher voltages. Such metal oxide transistors also allow for less leakage in the "off" state of the thin-film transistor (TFT) than can be achieved by, e.g., amorphous silicon TFTs. In a typical scanning TFT backplane comprising n lines, the transistor will be in the "off" state for approximately a proportion $(n-1)/n$ of the time required to refresh every line of the display. Any leakage of charge from the storage capacitors associated with each pixel would result in degradation of the electro-optical performance of the display. TFTs typically include a gate electrode, a gate-insulating film (typically SiO_2), a metal source electrode, a metal drain electrode, and a metal oxide semiconductor film over the gate-insulating film, at least partially overlapping the gate electrode, source electrode, and drain electrode. Such backplanes are available from manufacturers such as Sharp/Foxconn, LG, and BOE. Such backplanes are able to provide driving voltages of $\pm 30\text{V}$ (or more). Intermediate voltage drivers can be included so that the resulting driving waveforms may include five levels, or seven levels, or nine levels, or more.

[0063] One preferred metal oxide material for such applications is indium gallium zinc oxide (IGZO). IGZO-TFT has 20–50 times the electron mobility of amorphous silicon. By using IGZO TFTs in an active matrix backplane, it is possible to provide voltages of greater than 30V via a suitable display driver. This enables use of a source driver (a switch in the EPD that determines which voltage is applied to each of the column electrodes for a given selected row of the display) capable of supplying at least five, and perhaps seven driving voltage levels. In one example, there are two positive voltages, two negative voltages, and zero volts. In another example, there are three positive voltages, three negative voltages, and zero volts. In another example, there are four positive voltages, four negative voltages, and zero volts. These levels may be chosen within the range of about -27V to +27V, without the limitations imposed by top plane switching as described above.

[0064] The electrophoretic medium shown in FIGS. 2A-2E includes four types of electrophoretic particles in a non-polar fluid 17. A first particle (W-*; open circle) is negatively charged and may be surface treated so that the electrophoretic mobility of the first particle is dependent upon the strength of the driving electric field (discussed in greater detail below). In such instances, the electrophoretic mobility of the particle actually decreases in the presence of a stronger electric field, which is somewhat counter-intuitive. A second particle (M++*; dark circle) is positively charged, and may also be surface treated (or purposely untreated) so that either the electrophoretic mobility of the second particle is dependent upon the strength of the driving electric field, or the rate of unpacking of a collection of the second particle, after having been driven to one side of the cavity containing the particles upon reversal of the electric field direction, is slower than the rate of unpacking of collections of the third and fourth particles, or the particle forms a Coulombic aggregate with the first particle (W- in this case) that is separable by a high applied electric field, but not by a low applied electric field. A third particle (Y+; checkered circle) is positive, but has a charge magnitude that is smaller than the second particle. Additionally, the third particle may be surface treated, but not in a way that causes the electrophoretic mobility of the third particle to depend upon the strength of the driving electric field. That is, the third particle may have a surface treatment, however such a surface treatment does not result in the aforementioned reduction in electrophoretic mobility with an increased electric field. The fourth particle (C+++; gray circle) has the highest magnitude positive charge and the same type of surface treatment as the third particle. As indicated in FIG. 2A, the particles are nominally white, magenta, yellow, and cyan in color to produce colors as shown in FIG. 1. However, the system is not limited to this specific color set, nor is it limited to one

reflective particle and three absorptive particles. For example, the system could include one black absorptive particle and three reflective particles of red, yellow, and blue with suitably matched reflectance spectra to produce a process white state when all three reflective particles are mixed and viewable at the surface.

[0065] The first particle (negative) is white and scattering. The second particle (positive, medium charge magnitude) is magenta and absorptive. The third particle (positive, low charge magnitude) is yellow and absorptive. The fourth particle (positive, high charge magnitude) is cyan and absorptive. Table 1 below shows the diffuse reflectance of exemplary yellow, magenta, cyan and white particles useful in electrophoretic media of the present invention, together with the ratio of their absorption and scattering coefficients according to the Kubelka-Munk analysis of these materials as dispersed in a poly(isobutylene) matrix.

Table 1. Diffuse reflectance of preferred yellow, magenta, cyan and white particles.

		Diffuse reflectance of 1 μ m layer on 0% black			Ratio absorption / scatter		
Color	Volume Fraction	450 nm	550 nm	650 nm	K/S 450nm	K/S 550nm	K/S 650nm
Yellow (Y1)	0.097	4.5%	0.9%	0.5%	9.67	0.38	0.63
Yellow (Y1)	0.147	4.4%	0.9%	0.4%	9.84	0.25	0.02
Magenta (M1)	0.115	2.8%	3.8%	0.7%	10.01	10.85	1.27
Magenta (M1)	0.158	3.2%	4.1%	1.0%	10.00	10.75	1.64
Magenta (M1)	0.190	3.4%	4.1%	1.3%	10.09	10.80	1.03
Cyan (C1)	0.112	1.3%	3.7%	4.3%	7.27	11.17	10.22
Cyan (C1)	0.157	1.5%	3.8%	4.3%	7.41	11.30	10.37
Cyan (C1)	0.202	1.7%	3.9%	4.3%	7.21	11.56	10.47
White (W1)	0.147	8.1%	6.2%	4.8%	0.0015	0.0020	0.0026
White (W1)	0.279	24.9%	20.6%	17.0%	0.0003	0.0003	0.0004
White (W1)	0.339	26.3%	21.7%	18.1%	0.0001	0.0002	0.0002

[0066] The electrophoretic medium may be in any of the forms discussed above. Thus, the electrophoretic medium may be unencapsulated, encapsulated in discrete capsules surrounded

by capsule walls, encapsulated in sealed microcells, or in the form of a polymer-dispersed medium. The pigments are described in further detail elsewhere, such as in U.S. Patent Nos. 9,697,778 and 9,921,451. Briefly, white particle W1 is a silanol-functionalized light-scattering pigment (titanium dioxide) to which a polymeric material comprising lauryl methacrylate (LMA) monomers has been attached as described in U.S. Patent No. 7,002,728. White particle W2 is a polymer-coated titania produced substantially as described in Example 1 of U.S. Patent No. 5,852,196, with a polymer coating comprising an approximately 99:1 ratio of lauryl methacrylate and 2,2,2-trifluoroethyl methacrylate. Yellow particle Y1 is C.I. Pigment Yellow 180, used without coating and dispersed by attrition in the presence of Solsperse 19000, as described generally in U.S. Patent No. 9,697,778. Yellow particle Y2 is C.I. Pigment Yellow 155 used without coating and dispersed by attrition in the presence of Solsperse 19000, as described generally in U.S. Patent No. 9,697,778. Yellow particle Y3 is C.I. Pigment Yellow 139, used without coating and dispersed by attrition in the presence of Solsperse 19000, as described generally in U.S. Patent No. 9,697,778. Yellow particle Y4 is C.I. Pigment Yellow 139, which is coated by dispersion polymerization, incorporating trifluoroethyl methacrylate, methyl methacrylate and dimethylsiloxane-containing monomers as described in Example 4 of U.S. Patent No. 9,921,451. Magenta particle M1 is a positively-charged magenta material (dimethylquinacridone, C.I. Pigment Red 122) coated using vinylbenzyl chloride and LMA as described in U.S. Patent No. 9,697,778 and in Example 5 of U.S. Patent No. 9,921,451.

[0067] Magenta particle M2 is a C.I. Pigment Red 122, which is coated by dispersion polymerization, methyl methacrylate and dimethylsiloxane-containing monomers as described in Example 6 of U.S. Patent No. 9,921,451. Cyan particle C1 is a copper phthalocyanine material (C.I. Pigment Blue 15:3) which is coated by dispersion polymerization, incorporating methyl methacrylate and dimethylsiloxane-containing monomers as described in Example 7 of U.S. Patent No. 9,921,451. In some embodiments, it has been found that the color gamut is improved by using Ink Jet Yellow 4GC (Clariant) as the core yellow pigment, with incorporation of methyl methacrylate surface polymers. The zeta potential of this yellow pigment can be tuned with the addition of 2,2,2-trifluoroethyl methacrylate (TFEM) monomers and monomethacrylate terminated poly(dimethylsiloxane).

[0068] Electrophoretic media additives and surface treatments for facilitating differential electrophoretic mobility, as well as proposed mechanisms for interaction between the surface treatment and surrounding charge control agents and/or free polymers, are discussed in detail in U.S. Patent No. 9,697,778, incorporated by reference in its entirety. In such electrophoretic

media, one way of controlling the interactions among the various types of particles is by controlling the kind, amount, and thickness of polymeric coatings on the particles. For example, to control the particle characteristics such that the particle-particle interactions are less between the second type of particles and the third and fourth types of particles than between, e.g., the third type of particles and the fourth type of particles of the third species, the second type of particle may bear a polymeric surface treatment, while the third and fourth types of particles bear either no polymeric surface treatment or a polymeric surface treatment having a lower mass coverage per unit area of the particle surface than the second type of particles. More generally, the Hamaker constant (which is a measure of the strength of the Van der Waals interaction between two particles, the pair potential being proportional to the Hamaker constant and inversely proportional to the sixth power of the distance between the two particles) and/or the interparticle spacing need(s) to be adjusted by judicious choice of the polymeric coating(s) on the three species of particles.

[0069] As discussed in U.S. Patent No. 9,921,451, different types of polymers may include different types of polymer surface treatment. For example, Coulombic interactions may be weakened when the closest distance of approach of oppositely-charged particles is maximized by a steric barrier (typically a polymer grafted or adsorbed to the surface of one or both particles). The polymer shell may be a covalently-bonded polymer made by grafting processes or chemisorption as is well known in the art, or may be physisorbed onto the particle surface. For example, the polymer may be a block copolymer comprising insoluble and soluble segments. Alternatively, the polymer shell may be dynamic in that it is a loose network of free polymer from the electrophoretic medium that is complexed with a pigment particle in the presence of an electric field and a sufficient amount and kind of charge control agent (CCA – discussed below). Thus, depending upon the strength and polarity of the electric field, a particle may have more associated polymer, which causes the particle to interact differently with the container (e.g., microcapsule or microcell) and the other particles. The extent of the polymer shell is conveniently assessed by thermal gravimetric analysis (TGA), a technique in which the temperature of a dried sample of the particles is raised and the mass loss due to pyrolysis is measured as a function of temperature. Using TGA, the proportion of the mass of the particle that is polymer can be measured, and this can be converted to a volume fraction using the known densities of the core pigments and the polymers attached to them. Conditions can be found in which the polymer coating is lost but the core pigment remains (these conditions depend upon the precise core pigment particle used). A variety of polymer combinations can

be made to work as described below with respect to FIGS. 2A-2E. For example, in some embodiments a particle (typically the first and/or second particle) can have a covalently-attached polymer shell that interacts strongly with the container (e.g., microcell or microcapsule). Meanwhile the other particles of the same charge have no polymer coating or complex with free polymers in the solution so that those particles have little interaction with the container. In other embodiments, a particle (typically the first and/or second particle) will have no surface coating so that it is easier for that particle to form a charge double layer and experience electrophoretic mobility reduction in the presence of strong fields.

[0070] The fluid 17, in which the four types of particles are dispersed, is clear and colorless. The fluid contains the charged electrophoretic particles, which move through the fluid under the influence of an electric field. A preferred suspending fluid has a low dielectric constant (about 2), high volume resistivity (about 10^{15} Ohm-cm), low viscosity (less than 5 mPas), low toxicity and environmental impact, low water solubility (less than 10 parts per million (ppm), if traditional aqueous methods of encapsulation are to be used; note however that this requirement may be relaxed for non-encapsulated or certain microcell displays), a high boiling point (greater than about 90°C), and a low refractive index (less than 1.5). The last requirement arises from the use of scattering (typically white) pigments of high refractive index, whose scattering efficiency depends upon a mismatch in refractive index between the particles and the fluid.

[0071] Organic solvents such as saturated linear or branched hydrocarbons, silicone oils, halogenated organic solvents, and low molecular weight halogen-containing polymers are some useful fluids. The fluid may comprise a single component or may be a blend of more than one component in order to tune its chemical and physical properties. Reactants or solvents for the microencapsulation process (if used), such as oil soluble monomers, can also be contained in the fluid.

[0072] The fluid 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 fluids include hydrocarbons such as Isopar®, decahydronaphthalene (DECALIN), 5-ethylidene-2-norbornene, fatty oils, paraffin oil, silicon fluids, aromatic hydrocarbons such as toluene, xylene, phenylxylylethane, dodecylbenzene or alkylnaphthalene, halogenated solvents such as perfluorodecalin, perfluorotoluene, perfluoroxylene, dichlorobenzotrifluoride, 3,4,5 -trichlorobenzotri fluoride, chloropentafluorobenzene, dichlorononane or pentachlorobenzene, and perfluorinated solvents such as FC-43,

FC-70 or FC-5060 from 3M Company, St. Paul MN, low molecular weight halogen containing polymers such as poly(perfluoropropylene oxide) from TCI America, Portland, Oregon, poly(chlorotrifluoro-ethylene) such as Halocarbon Oils from Halocarbon Product Corp., River Edge, NJ, perfluoropolyalkylether such as Galden from Ausimont or Krytox Oils and Greases K-Fluid Series from DuPont, Delaware, polydimethylsiloxane based silicone oil from Dow-corning (DC -200).

[0073] The electrophoretic media typically also include one or more charge control agents (CCA), and may also include a charge director. CCA and charge directors typically comprise low molecular weight surfactants, polymeric agents, or blends of one or more components and serve to stabilize or otherwise modify the sign and/or magnitude of the charge on the electrophoretic particles. The CCA is typically a molecule comprising ionic or other polar groupings, hereinafter referred to as head groups. At least one of the positive or negative ionic head groups is preferably attached to a non-polar chain (typically a hydrocarbon chain) that is hereinafter referred to as a tail group. It is thought that the CCA forms reverse micelles in the internal phase and that it is a small population of charged reverse micelles that leads to electrical conductivity in the very non-polar fluids typically used as electrophoretic fluids.

[0074] The addition of CCAs provides for the production of reverse micelles including a highly polar core that may vary in size from 1 nm to tens of nanometers (and may have spherical, cylindrical, or other geometry) surrounded by the non-polar tail groups of the CCA molecule. In electrophoretic media, three phases may typically be distinguished: a solid particle having a surface, a highly polar phase that is distributed in the form of extremely small droplets (reverse micelles), and a continuous phase that comprises the fluid. Both the charged particles and the charged reverse micelles may move through the fluid upon application of an electric field, and thus there are two parallel pathways for electrical conduction through the fluid (which typically has a vanishingly small electrical conductivity itself).

[0075] The polar core of the CCA is thought to affect the charge on surfaces by adsorption onto the surfaces. In an EPD, such adsorption may be onto the surfaces of the electrophoretic particles or the interior walls of a microcapsule (or other solid phase, such as the walls of a microcell) to form structures similar to reverse micelles, these structures hereinafter being referred to as hemi-micelles. When one ion of an ion pair is attached more strongly to the surface than the other (e.g., by covalent bonding), ion exchange between hemi-micelles and unbound reverse micelles can lead to charge separation in which the more strongly bound ion

remains associated with the particle and the less strongly bound ion becomes incorporated into the core of a free reverse micelle.

[0076] It is also possible that the ionic materials forming the head group of the CCA may induce ion-pair formation at the particle (or other) surface. Thus the CCA may perform two basic functions: charge-generation at the surface and charge-separation from the surface. The charge-generation may result from an acid-base or an ion-exchange reaction between some moiety present in the CCA molecule or otherwise incorporated into the reverse micelle core or fluid, and the particle surface. Thus, useful CCA materials are those which are capable of participating in such a reaction, or any other charging reaction as known in the art.

[0077] Non-limiting classes of charge control agents that are useful in electrophoretic media include organic sulfates or sulfonates, metal soaps, block or comb copolymers, organic amides, organic zwitterions, and organic phosphates and phosphonates. Useful organic sulfates and sulfonates include, but are not limited to, sodium bis(2-ethylhexyl) sulfosuccinate, calcium dodecylbenzenesulfonate, calcium petroleum sulfonate, neutral or basic barium dinonylnaphthalene sulfonate, neutral or basic calcium dinonylnaphthalene sulfonate, dodecylbenzenesulfonic acid sodium salt, and ammonium lauryl sulfate. Useful metal soaps include, but are not limited to, basic or neutral barium petronate, calcium petronate, cobalt, calcium, copper, manganese, magnesium, nickel, zinc, aluminum and iron salts of carboxylic acids such as naphthenic, octanoic, oleic, palmitic, stearic, and myristic acids and the like. Useful block or comb copolymers include, but are not limited to, AB diblock copolymers of (A) polymers of 2-(N,N-dimethylamino)ethyl methacrylate quaternized with methyl p-toluenesulfonate and (B) poly(2-ethylhexyl methacrylate), and comb graft copolymers with oil soluble tails of poly(12-hydroxystearic acid) and having a molecular weight of about 1800, pendant on an oil-soluble anchor group of poly(methyl methacrylate-methacrylic acid). Useful organic amides/amines include, but are not limited to, polyisobutylene succinimides such as OLOA 371 or 1200 (available from Chevron Oronite Company LLC, Houston, Tex.), or SOLSPERSE 17000 or 19000 (available from Lubrizol, Wickliffe, OH: Solsperse is a Registered Trade Mark), and N-vinylpyrrolidone polymers. Useful organic zwitterions include, but are not limited to, lecithin. Useful organic phosphates and phosphonates include, but are not limited to, the sodium salts of phosphated mono- and di-glycerides with saturated and unsaturated acid substituents. Useful tail groups for CCA include polymers of olefins such as poly(isobutylene) of molecular weight in the range of 200 – 10,000. The head groups may be sulfonic, phosphoric or carboxylic acids or amides, or alternatively amino groups such as

primary, secondary, tertiary or quaternary ammonium groups. One class of CCAs that are useful in the disclosed four-particle electrophoretic media are disclosed in U.S. Patent Publication No. 2017/0097556, incorporated by reference herein in its entirety. Such CCAs typically include a quaternary amine head group and an unsaturated polymeric tail, i.e., including at least one C-C double bond. The polymeric tail is typically a fatty acid tail. A variety of CCA molecular weights can be used. In some embodiments, the molecular weight of the CCA is 12,000 grams/mole or greater, e.g., between 14,000 grams/mole and 22,000 grams/mole.

[0078] Charge adjuvants used in the media may bias the charge on electrophoretic particle surfaces, as described in more detail below. Such charge adjuvants may be Bronsted or Lewis acids or bases. Exemplary charge adjuvants are disclosed in U.S. Patent Nos. 9,765,015, 10,233,339 and 10,782,586, all of which are incorporated by reference in their entireties. Exemplary adjuvants may include polyhydroxy compounds which contain at least two hydroxyl groups include, but are not limited to, ethylene glycol, 2,4,7,9-tetramethyldecyne-4,7-diol, poly(propylene glycol), pentaethylene glycol, tripropylene glycol, triethylene glycol, glycerol, pentaerythritol, glycerol tris(12-hydroxystearate), propylene glycerol monohydroxystearate, and ethylene glycol monohydroxystearate. Examples of aminoalcohol compounds which contain at least one alcohol function and one amine function in the same molecule include, but are not limited to, triisopropanolamine, triethanolamine, ethanolamine, 3-amino-1-propanol, o-aminophenol, 5-amino-1-pentanol, and tetrakis(2-hydroxyethyl)ethylenediamine. In some embodiments, the charge adjuvant is present in the EPD medium in an amount of between about 1 to about 500 milligrams per gram ("mg/g") of the particle mass, and more preferably between about 50 to about 200 mg/g.

[0079] Particle dispersion stabilizers may be added to prevent particle flocculation or attachment to the capsule or other walls or surfaces. For the typical high resistivity liquids used as fluids in EPDs, non-aqueous surfactants may be used. These include, but are not limited to, glycol ethers, acetylenic glycols, alkanolamides, sorbitol derivatives, alkyl amines, quaternary amines, imidazolines, dialkyl oxides, and sulfosuccinates.

[0080] As described in U.S. Patent No. 7,170,670, the bistability of electrophoretic media can be improved by including in the fluid a polymer having a number average molecular weight in excess of about 20,000, this polymer being essentially non-absorbing on the electrophoretic particles; poly(isobutylene) is a preferred polymer for this purpose. Also, as described in e.g., U.S. Patent No. 6,693,620, a particle with immobilized charge on its surface sets up an electrical double layer of opposite charge in a surrounding fluid. Ionic head groups of the CCA

may be ion-paired with charged groups on the electrophoretic particle surface, forming a layer of immobilized or partially immobilized charged species. Outside this layer is a diffuse layer comprising charged (reverse) micelles comprising CCA molecules in the fluid. In conventional DC electrophoresis an applied electric field exerts a force on the fixed surface charges and an opposite force on the mobile counter-charges, such that slippage occurs within the diffuse layer and the particle moves relative to the fluid. The electric potential at the slip plane is known as the zeta potential.

[0081] As a result, some of the particle types within the electrophoretic medium have different electrophoretic mobilities depending upon the strength of the electric field across the electrophoretic medium. For example, when a first (low strength, i.e., around $\pm 10\text{V}$ or less) electric field is applied to the electrophoretic medium, the first type of particles move in one direction relative to the electric field, however, when a second (high strength, i.e., around $\pm 20\text{V}$ or more) electric field is applied, having the same polarity as the first electric field, the first type of particles begins to move in the opposite direction relative to the electric field. It is theorized that the behavior results from conduction within the highly non-polar fluid being mediated by charged reverse micelles or counter-charged electrophoretic particles. Accordingly, any electrochemically-generated protons (or other ions) are probably transported through the non-polar fluid in micelle cores or adsorbed on electrophoretic particles. For example, as illustrated in FIG. 5B of U.S. Patent No. 9,697,778, a positively-charged reverse micelle may approach a negative electrophoretic particle traveling in the opposite direction, wherein the reverse micelle is incorporated into the electric double layer around the negatively charged particle. (The electric double layer includes both the diffuse layer of charge with enhanced counter-ion concentration and the hemi-micellar surface-adsorbed coating on the particle; in the latter case, the reverse micelle charge would become associated with the particle within the slip envelope that, as noted above, defines the zeta potential of the particle.) Through this mechanism, an electrochemical current of positively-charged ions flows through the electrophoretic fluid, and the negatively-charged particles may become biased towards a more positive charge. As a result, the electrophoretic mobility, e.g., of the first negative type of particle is a function of the magnitude of the electrochemical current and the residence time of a positive charge close to the particle surface, which is a function of the strength of the electric field.

[0082] Furthermore, as also described in U.S. Patent No. 9,697,778, positively-charged particles can be prepared that also exhibit different electrophoretic mobilities depending upon

the applied electric field. In some embodiments, a secondary (or co-) CCA can be added to the electrophoretic medium to adjust the zeta potentials of the various particles. Careful selection of the co-CCA may allow alteration of the zeta potential of one particle while leaving those of the other particles essentially unchanged, allows close control of both the electrophoretic velocities of the various particles during switching and the inter-particle interactions.

[0083] In some embodiments, a portion of the charge control agents intended for the final formulation are added during synthesis of the electrophoretic particles to engineer the desired zeta potential and to influence the reduction in electrophoretic mobility due to a strong electric field. For example, it has been observed that adding quaternary amine charge control agents during polymer grafting will result in some amount of the CCA being complexed to the particles. (This can be confirmed by removing the particles from the electrophoretic fluid and subsequently stripping the surface species from the pigments with THF to remove all adsorbed species. When the THF extract is evaluated with ¹H NMR, it is clear that a good amount of the CCA was adsorbed to the pigment particles or complexed with the surface polymer.) Experiments suggest that high CCA loading amongst the surface polymers of the particles facilitates the formation of a charge double layer around the particle in the presence of a strong electric field. For example, magenta particles having greater than 200 mg of a charge control agent (CCA) per gram of finished magenta particle have excellent staying properties in the presence of a high positive electric field. (See, e.g., FIG. 2C, and the description above.) In some embodiments, the CCA includes a quaternary amine head group and a fatty acid tail. In some embodiments the fatty acid tail is unsaturated. When some of the particles in the electrophoretic medium include high CCA loading, it is important that the particles for which consistent electrophoretic mobility is desired do not have substantial CCA loading, e.g., less than 50 mg of a charge control agent (CCA) per gram of finished particle, e.g., less than 10 mg of a charge control agent (CCA) per gram of finished particle.

[0084] Alternately, an electrophoretic medium including four types of particles in the presence of Solspers 17000 in Isopar E benefits from the additions of small amounts of acidic entities, such as, e.g., aluminum salts of di-*t*-butyl salicylic acid (Bontron E-88, available from Orient Corporation, Kenilworth, NJ)). Addition of the acidic material moves the zeta potential of many particles (though not all) to more positive values. In one case, about 1% of the acidic material and 99% of Solspers 17000 (based on total weight of the two materials) moves the zeta potential of the third type of particle (Y+) from -5mV to about +20mV. Whether or not the

zeta potential of a particular particle is changed by a Lewis acidic material like the aluminum salt will depend upon the details of the surface chemistry of the particle.

[0085] Table 2 shows exemplary relative zeta potentials of the three types of colored and singular white particles in a preferred embodiment.

Table 2. Relative zeta potentials of colored particles in the presence of relative zeta potential of white particles.

		White zeta potential (mV)			
		-30	0	10	20
Cyan zeta potential (mV)	80	110	80	70	60
Magenta zeta potential (mV)	40	70	40	30	20
+Yellow zeta potential (mV)	20	50	20	10	0
-Yellow zeta potential (mV)	-20	10	-20	-30	-40

[0086] The negative (white) particle has a zeta potential of -30mV, and the remaining three particles are all positive relative to the white particle. Accordingly, a display comprising positive cyan, magenta, and yellow particles can switch between a black state (with all colored particles in front of the white particle with respect to the viewing surface) and a white state, with the white particle closest to the viewer, and blocking the viewer from perceiving the remaining three particles. In contrast, when the white particle has a zeta potential of 0V, the negatively-charged yellow particle is the most negative of all the particles, and thus a display comprising this particle would switch between a yellow and a blue state. This would also occur if the white particle were positively charged. The positively-charged yellow particle, however, would be more positive than the white particle unless its zeta potential exceeded +20mV.

[0087] The behavior of the electrophoretic media is consistent with the mobility of the white particle (represented in Table 2 as the zeta potential) being dependent upon the applied electric field. Thus, in the example illustrated in Table 2, when addressed with a low voltage the white particle might behave as though its zeta potential were -30mV, but when addressed with a higher voltage it might behave as though its zeta potential were more positive, maybe even as high as +20mV (matching the zeta potential of the yellow particle). Thus, when addressed with a low voltage, the display would switch between black and white states, but when addressed at a higher voltage would switch between blue and yellow states.

[0088] It is also possible that particles of opposite charge may form Coulombic aggregates. In the aggregated form the mobility of the particles may be different from what would be measured for the components of the aggregate. Thus, for example, an aggregate may be formed between the negatively-charged white particles and any of the oppositely-charge pigments. In

certain embodiments it may be preferred that the electric field required to separate Coulombic aggregates be higher for some pigment pairs than for others. In the present case it is likely that the electric field required to separate an aggregate formed between the magenta and white pigments is larger than that required to separate aggregates formed between the cyan and white or yellow and white particles.

[0089] It is possible that the charges of the various pigment particles are affected by the presence of other charged pigments in their environment. Thus, it is not necessarily true that the charges (or zeta potentials) of the pigments in their final formulations are the same as those measured for the pigments on their own as dispersed in a solvent in the presence of a CCA.

[0090] The motion of the various particles according to one possible hypothesis in the presence of a high (e.g., “ $\pm H$ ”, e.g., $\pm 20V$, e.g., $\pm 25V$) electric field and a low (e.g., “ $\pm L$ ”, e.g., $\pm 5V$, e.g., $\pm 10V$) electric field are shown in FIGS. 2B-2E. For the purposes of illustration, each box bounded by dashed lines represents a pixel bounded by a top light-transmissive electrode 21 and a bottom electrode 22, which may be a pixel electrode of an active matrix, however it may also be a light-transmissive electrode, or a segmented electrode, etc. Starting from a first state, in which all of the positive particles are present at the viewing surface (nominally black), the electrophoretic medium can be driven to four different optical states, as shown in FIGS. 2B-2E. In the preferred embodiment, this results in a white optical state (FIG. 2B), a magenta optical state (FIG. 2C), a yellow optical state (FIG. 2D), and a red optical state (FIG. 2E). It should be evident that the remaining four optical states of FIG. 1 can be achieved by reversing the order of the initial state and the driving electric fields, as shown in short hand in FIG. 5.

[0091] When addressed with a low voltage, as in FIG. 2B, the particles behave according to their relative zeta potentials with relative velocities illustrated by the arrows for the case when a negative voltage is applied to the backplane. Thus, in this example, the cyan particles move faster than the magenta particles, which move faster than the yellow particles. The first (positive) pulse does not change the positions of the particles, since they are already restricted in motion by the walls of the enclosure. The second (negative) pulse exchanges the positions of the colored and white particles, and thus the display switches between black and white states, albeit with transient colors reflecting the relative mobilities of the colored particles. Reversing the starting positions and polarities of the pulses allows for a transition from white to black. Accordingly, this embodiment provides black-white updates that require lower voltages (and consume less power) as compared to other black and white formulations achieved with multiple colors via either a process black or a process white.

[0092] In FIG. 2C, the first (positive) pulse is of a high positive voltage, sufficient to reduce the mobility of the magenta particle (i.e., the particle of intermediate mobility of the three positively-charge colored particles). Because of the reduced mobility, the magenta particles essentially remain frozen in place, and a subsequent pulse in the opposite direction, of low voltage, moves the cyan, white and yellow particles more than the magenta particles, thereby producing a magenta color at the viewing surface, with the negative white particles behind the magenta particles. Importantly, if the starting position and the polarities of the pulses are reversed, (equivalent to viewing the display from the side opposite the viewing surface, i.e., through electrode 22), this pulse sequence would produce a green color (i.e., a mixture of yellow and cyan particles).

[0093] In FIG. 2D, the first pulse is of a low voltage that does not significantly reduce the mobility of the magenta particles or the white particles. However, the second pulse is of a high negative voltage that reduces the mobility of the white particles. This allows more effective racing between the three positive particles, such that the slowest type of particles (yellow in this example) remains visible in front of the white particle, whose movement was diminished with the earlier negative pulse. Notably, the yellow particles do not make it to the top surface of the cavity containing the particles. Importantly, if the starting position and the polarities of the pulses are reversed, (equivalent to viewing the display from the side opposite the viewing surface, i.e., through electrode 22), this pulse sequence would produce a blue color (i.e., a mixture of magenta and cyan particles).

[0094] FIG. 2E shows that when both pulses are of high voltage, the magenta particle mobility would be reduced by the first high positive pulse, and the racing between cyan and yellow would be enhanced by the reduction in white mobility caused by the second high negative pulse. This produces a red color. Importantly, if the starting position and the polarities of the pulses are reversed, (equivalent to viewing the display from the side opposite the viewing surface, i.e., through electrode 22), this pulse sequence would produce a cyan color.

[0095] To obtain a high-resolution display, individual pixels of a display should be addressable without interference from adjacent pixels. One way to achieve this objective is to provide an array of non-linear elements, such as transistors or diodes, with at least one non-linear element associated with each pixel, to produce an "active matrix" display. An addressing or pixel electrode, which addresses one pixel, is connected to an appropriate voltage source through the associated non-linear element. Typically, when the non-linear element is a transistor, the pixel electrode is connected to the drain of the transistor, and this arrangement

will be assumed in the following description, although it is essentially arbitrary and the pixel electrode could be connected to the source of the transistor. Conventionally, in high resolution arrays, the pixels are arranged in a two-dimensional array of rows and columns, such that any specific pixel is uniquely defined by the intersection of one specified row and one specified column. The sources of all the transistors in each column are connected to a single column electrode, while the gates of all the transistors in each row are connected to a single row electrode; again the assignment of sources to rows and gates to columns is conventional but essentially arbitrary, and could be reversed if desired. The row electrodes are connected to a row driver, which essentially ensures that at any given moment only one row is selected, i.e., that there is applied to the selected row electrode a select voltage such as to ensure that all the transistors in the selected row are conductive, while there is applied to all other rows a non-select voltage such as to ensure that all the transistors in these non-selected rows remain non-conductive. The column electrodes are connected to column drivers, which place upon the various column electrodes voltages selected to drive the pixels in the selected row to their desired optical states. (The aforementioned voltages are relative to a common front electrode, which is conventionally provided on the opposed side of the electro-optic medium from the non-linear array and extends across the whole display.) After a pre-selected interval known as the "line address time" the selected row is deselected, the next row is selected, and the voltages on the column drivers are changed so that the next line of the display is written. This process is repeated so that the entire display is written in a row-by-row manner.

[0096] Conventionally, each pixel electrode has associated therewith a capacitor electrode such that the pixel electrode and the capacitor electrode form a capacitor; see, e.g., International Patent Application WO2001007961. In some embodiments, N-type semiconductor (e.g., amorphous silicon) may be used to form the transistors and the "select" and "non-select" voltages applied to the gate electrodes can be positive and negative, respectively.

[0097] FIG. 3 of the accompanying drawings depicts an exemplary equivalent circuit of a single pixel of an EPD. As illustrated, the circuit includes a capacitor 10 formed between a pixel electrode and a capacitor electrode. The electrophoretic medium 20 is represented as a capacitor and a resistor in parallel. In some instances, direct or indirect coupling capacitance 30 between the gate electrode of the transistor associated with the pixel and the pixel electrode (usually referred to as a "parasitic capacitance") may create unwanted noise to the display. Usually, the parasitic capacitance 30 is much smaller than that of the storage capacitor 10, and when the pixel rows of a display is being selected or deselected, the parasitic capacitance 30

may result in a small negative offset voltage to the pixel electrode, also known as a “kickback voltage”, which is usually less than 2 volts. In some embodiments, to compensate for the unwanted “kickback voltage”, a common potential V_{com} , may be supplied to the top plane electrode and the capacitor electrode associated with each pixel, such that, when V_{com} is set to a value equal to the kickback voltage (V_{KB}), every voltage supplied to the display may be offset by the same amount, and no net DC-imbalance experienced.

[0098] Problems may arise, however, when V_{com} is set to a voltage that is not compensated for the kickback voltage. This may occur when it is desired to apply a higher voltage to the display than is available from the backplane alone. It is well known in the art that, e.g., the maximum voltage applied to the display may be doubled if the backplane is supplied with a choice of a nominal $+V$, 0 , or $-V$, e.g., while V_{com} is supplied with $-V$. The maximum voltage experienced in this case is $+2V$ (i.e., at the backplane relative to the top plane), while the minimum is zero. If negative voltages are needed, the V_{com} potential must be raised at least to zero. Waveforms used to address a display with positive and negative voltages using top plane switching must therefore have particular frames allocated to each of more than one V_{com} voltage setting.

[0099] A set of waveforms for driving a color EPD having four particles described in U.S. Patent No. 9,921,451, which is incorporated by reference herein. In U.S. Patent No. 9,921,451, seven different voltages are applied to the pixel electrodes: three positive, three negative, and zero. However, in some cases, the maximum voltages used in these waveforms are higher than that can be handled by amorphous silicon thin-film transistors. In such instances, suitable high voltages can be obtained by the use of top plane switching. When (as described above) V_{com} is deliberately set to V_{KB} , a separate power supply may be used. It is costly and inconvenient, however, to use as many separate power supplies as there are V_{com} settings when top plane switching is used. Furthermore, top plane switching is known to increase kickback, thereby degrading the stability of the color states.

[0100] A display device may be constructed using an electrophoretic fluid in several ways that are known in the prior art. The electrophoretic fluid may be encapsulated in microcapsules or incorporated into microcell structures that are thereafter sealed with a polymeric layer. The microcapsule or microcell layers may be coated or embossed onto a plastic substrate or film bearing a transparent coating of an electrically conductive material. This assembly may be laminated to a backplane bearing pixel electrodes using an electrically conductive adhesive. Alternatively, the electrophoretic fluid may be dispensed directly on a thin open-cell grid that

has been arranged on a backplane including an active matrix of pixel electrodes. The filled grid can then be top-sealed with an integrated protective sheet/light-transmissive electrode.

[0101] FIG. 4 shows a schematic, cross-sectional drawing (not to scale) of a display structure 200 including an electrophoretic medium. In display 200, the electrophoretic fluid is illustrated as being confined to microcells, although other structures, e.g., microcapsules may also be used. Substrate 202, which may be glass or plastic, bears pixel electrodes 204 that are either individually addressed segments or associated with thin film transistors in an active matrix arrangement. (The combination of substrate 202 and electrodes 204 is conventionally referred to as the back plane of the display.) Layer 206 is an optional dielectric layer according to the invention applied to the backplane. (Methods for depositing a suitable dielectric layer are described in U.S. Patent Publication No. 2020/0348576, incorporated by reference.) The front plane of the display comprises transparent substrate 222 that bears a transparent, electrically conductive coating 220. Overlying electrode layer 220 is an optional dielectric layer 218. Layer (or layers) 216 are polymeric layer(s) that may comprise a primer layer for adhesion of microcells to transparent electrode layer 220 and some residual polymer comprising the bottom of the microcells. The walls of the microcells 212 are used to contain the electrophoretic fluid 214. The microcells are sealed with layer 210 and the whole front plane structure is adhered to the backplane using electrically-conductive adhesive layer 208. Processes for forming the microcells are described in the prior art, e.g., in U.S. Patent No. 6,930,818. In some instance, the microcells are less than 20 μm in depth, e.g., less than 15 μm in depth, e.g., less than 12 μm in depth, e.g., about 10 μm in depth, e.g., about 8 μm in depth.

[0102] Most commercial EPDs use amorphous silicon based thin-film transistors (TFTs) in the construction of active matrix backplanes (202/024) because of the wider availability of fabrication facilities and the costs of the various starting materials. Unfortunately, amorphous silicon thin-film transistors become unstable when supplied gate voltages that would allow switching of voltages higher than about $\pm 15\text{V}$. Nonetheless, as described below, the performance of ACeP is improved when the magnitudes of the high positive and negative voltages are allowed to exceed $\pm 15\text{V}$. Accordingly, as described in previous disclosures, improved performance is achieved by additionally changing the bias of the top light-transmissive electrode with respect to the bias on the backplane pixel electrodes, also known as top-plane switching. Thus, if a voltage of +30V (relative to the backplane) is needed, the top plane may be switched to -15V while the appropriate backplane pixel is switched to +15V.

Methods for driving a four-particle electrophoretic system with top-plane switching are described in greater detail in, e.g., U.S. Patent No. 9,921,451.

[0103] These waveforms require that each pixel of the display can be driven at five different addressing voltages, designated $+V_{\text{high}}$, $+V_{\text{low}}$, 0, $-V_{\text{low}}$ and $-V_{\text{high}}$, illustrated as 30V, 15V, 0, -15V and -30V. In practice, it may be preferred to use a larger number of addressing voltages. If only three voltages are available (i.e., $+V_{\text{high}}$, 0, and $-V_{\text{high}}$) it may be possible to achieve the same result as addressing at a lower voltage (say, V_{high}/n where n is a positive integer > 1) by addressing with pulses of voltage V_{high} but with a duty cycle of $1/n$.

[0104] FIG. 5 shows typical waveforms (in simplified form) used to drive the four-particle color electrophoretic display system described above. Such waveforms have a “push-pull” structure: i.e., they consist of a dipole comprising two pulses of opposite polarity. The magnitudes and lengths of these pulses determine the color obtained. The waveforms have five such voltage levels. FIG. 5 shows high and low positive and negative voltages, as well as zero volts. Typically, “low” (L) refers to a range of about 5 – 15V, while “high” (H) refers to a range of about 15 – 30V. In general, the higher the magnitude of the “high” voltages, the better the color gamut achieved by the display. An additional “medium” (M) level may also be used, which is typically around 15V; however, the value for M will depend somewhat on the composition of the particles, as well as the environment of the electrophoretic medium.

[0105] Although FIG. 5 shows the simplest dipoles required to form colors, it will be appreciated that practical waveforms may use multiple repetitions of these patterns, or other patterns that are aperiodic.

[0106] Of course, achieving the desired color with the driving pulses of FIG. 5 is contingent on the particles starting the process from a known state, which is unlikely to be the last color displayed on the pixel. Accordingly, a series of reset pulses precede the driving pulses, which increases the amount of time required to update a pixel from a first color to a second color. The reset pulses are described in greater detail in U.S. Patent No. 10,593,272, incorporated by reference. The lengths of these pulses (refresh and address) and of any rests (i.e., periods of zero voltage between them) may be chosen so that the entire waveform (i.e., the integral of voltage with respect to time over the whole waveform) is DC balanced (i.e., the integral of voltage over time is substantially zero). DC balance can be achieved by adjusting the lengths of the pulses and rests in the reset phase so that the net impulse supplied in the reset phase is equal in magnitude and opposite in sign to the net impulse supplied in the address phase, during which phase the display is switched to a particular desired color. As shown in the FIGS. 2B -

2E, however, the starting state for the eight primary colors is either a black or white state, which can be achieved with a sustained low voltage driving pulse. The simplicity of achieving this start state further reduces the time of updates between states, which is more pleasing for the user and also reduces the amount of power consumed (thus increasing battery life).

[0107] In addition, the foregoing discussion of the waveforms, and specifically the discussion of DC balance, ignores the question of kickback voltage. In practice, as previously, every backplane voltage is offset from the voltage supplied by the power supply by an amount equal to the kickback voltage V_{KB} . Thus, if the power supply used provides the three voltages $+V$, 0 , and $-V$, the backplane would actually receive voltages $V+V_{KB}$, V_{KB} , and $-V+V_{KB}$ (note that V_{KB} , in the case of amorphous silicon TFTs, is usually a negative number). The same power supply would, however, supply $+V$, 0 , and $-V$ to the front electrode without any kickback voltage offset. Therefore, e.g., when the front electrode is supplied with $-V$ the display would experience a maximum voltage of $2V+V_{KB}$ and a minimum of V_{KB} . Instead of using a separate power supply to supply V_{KB} to the front electrode, which can be costly and inconvenient, a waveform may be divided into sections where the front electrode is supplied with a positive voltage, a negative voltage, and V_{KB} .

FOUR PIGMENT EPD ADDRESSED BY FOUR VOLTAGE LEVELS

[0108] In accordance with one or more embodiments of the invention, a full-color EPD is disclosed that is addressable by waveforms having exactly four voltage levels. The EPD includes an electrophoretic medium having four types of charged electrophoretic pigment particles including three substantially non-light-scattering pigments that are colored, preferably corresponding to the subtractive primary colors (cyan, magenta, and yellow), and a fourth, light-scattering white pigment. The three colored pigments have one charge polarity, while the white pigment has the opposite charge polarity. The white pigment is preferably negatively charged, and the three colored pigments are positively charged. The display is addressed by waveforms having exactly four different voltage levels to render any of the primary colors (white, black, cyan, magenta, yellow, red, green, and blue) at any pixel location without utilizing top plane switching (i.e., while holding the voltage on the light-transmissive electrode of the EPD constant). The voltage levels are specified by two bits of data per pixel and preferably comprise high and low positive voltages, a single negative voltage, and a near zero voltage.

[0109] Because the EPD utilizes exactly four voltage levels and can be addressed by two bits of data per pixel, instead of three or more bits used in prior art color EPDs not using top-plane switching, device costs can be reduced while providing excellent performance and color quality without significant sacrifice of color gamut, as discussed below.

[0110] As noted above, U.S. Patent Nos. 9,697,778 and 10,678,111 disclose two different types of EPDs capable of rendering any color at any pixel location. The first of these patents discloses a display in which a white (light-scattering) pigment moves in a first direction when addressed with a low applied voltage and in the opposite direction when addressed with a higher voltage. The second patent discloses a full-color EPD containing four pigments: white, cyan, magenta, and yellow. U.S. Patent No. 10,276,109 and others disclose methods for addressing such displays using waveforms with at least five different voltage levels.

[0111] Prior art black and white EPDs typically use only three different voltage levels: one positive, one negative, and zero. The source driver of the black and white EPD accepts two bit data per pixel from the display controller, specifying which power supply is to be connected to which column.

[0112] When five voltage levels are needed (the minimum number used to address the full-color multi-pigment displays of the prior art), the module architecture must change in one of two general ways: (a) maintain the 2-bit per pixel architecture, but implement top-plane switching to change the potential difference between the backplane electrodes and the top plane (transparent) common electrode for every pixel in the display, or (b) provide a source driver capable of selecting between at least five different power supplies. In the latter case there is insufficient information in two bits, and at least three bits of data per pixel must be supplied by the controller to the source driver.

[0113] Both of these approaches have notable disadvantages. Since adjusting the top plane of the display affects every pixel, there is limited flexibility in the possible waveform structures that can be used to address the display. For example, it is not possible to assert the maximum positive and negative voltages across the electrophoretic fluid in two separate pixels at the same time. Consequently, the waveform is made longer, typically by a factor of about two. Requiring the controller to supply at least three bits of data per pixel requires faster data transfer rates (or longer transfer times), which either increases the cost of the controller or decreases the frame time that can be supported. In addition, many controllers in the current EPD ecosystem are incapable of supplying three bits of data per pixel. Providing at least five different voltage

levels can therefore lead to unacceptably high module costs or unduly long waveforms for many applications.

[0114] It would be desirable to address a color EPD using two bits of data per pixel without using top plane switching. Two bits of data per pixel can be used to specify four, rather than just three, voltage levels. Dynamic reassignment of power supplies on a per-frame basis would not provide an equally flexible alternative, as the reassignment would apply to every pixel when updating a particular frame and lead to the same type of compromise as top-plane switching.

[0115] In EPDs having a negatively-charged white pigment and positively-charged cyan, magenta, and yellow pigments, the complementary colors yellow/blue and magenta/green are best afforded by waveforms comprising dipoles (high- and low-voltage pulses of opposite polarity). Waveforms in prior art devices used approximately DC-balanced concatenated dipole sequences to render these colors. Thus, if a dipole comprises a first pulse of voltage V_1 for time t_1 , and a second pulse of voltage $-V_2$ for time t_2 , DC balance is achieved if $V_1 t_1 = V_2 t_2$.

[0116] When V_1 is positive and $|V_1| > |V_2|$, the display oscillates between the colors blue (at the end of the positive pulse) and magenta (at the end of the negative pulse). Conversely, if the polarities of each pulse are reversed, the display oscillates between the colors yellow (at the end of the negative pulse) and green (at the end of the positive pulse). Thus, it has been expected that all four of these voltage levels and a zero volt option would be needed to render all colors at each pixel.

[0117] However, it has been discovered that more complex, aperiodic voltage sequences having exactly four voltage levels (preferably high and low positive voltages, a single negative voltage, and a near zero voltage) can be used to address a display and make all the colors at every pixel location. (The term “aperiodic” refers to voltage sequences that do not have an exactly repeated structure.) Tuning such waveforms is more complex and difficult than the easily parameterized dipole structures of the prior art; so much so that it would require an impractically long time to perform the tuning directly with an actual display panel. However, it has been found that such complex waveforms can be more readily tuned using a two-stage computer modeling process. In the first stage, a set of arbitrary voltage lists is supplied to the display and the optical response is recorded at the end of every frame. From this training set, a model is constructed using machine learning methods that accurately predicts the effect of applying voltage lists that were not in the original training set (e.g., to an accuracy of about 1-

2 dE*). In the second stage, tuning of ideal waveforms is accomplished using the model, which allows vastly more waveforms to be tested in a given time than the physical display module.

[0118] FIGS. 6A and 6B show exemplary waveforms having four voltage levels for driving a four-particle color EPD in accordance with one or more embodiments. The figures show exemplary pulse sequences used to form the eight principal colors (red, green, blue, cyan, magenta, yellow, black, and white) at each pixel. The magnitudes and lengths of these pulses determine the color obtained. In one or more embodiments (e.g., as depicted in FIG. 6A), the four voltage levels comprise high and low positive voltages, near zero volts, and a high negative voltage (designated as $+V_{\text{high}}$, $+V_{\text{low}}$, 0, and $-V_{\text{high}}$). In one or more alternate embodiments (e.g., as depicted in FIG. 6B), the four voltage levels comprise high and low negative voltages, near zero volts, and a high positive voltage (designated as $-V_{\text{high}}$, $-V_{\text{low}}$, 0, and $+V_{\text{high}}$). Typically, the “low” voltage has a range of about 5 to 15V, while a “high” voltage has a range of about 15 to 30V. In general, the higher the magnitude of the “high” voltages, the better the color gamut achieved by the display. In one particular embodiment, the four voltage levels are 24V, 10V, 0, and -24V. In another particular embodiment, the four voltage levels are -24V, -10V, 0, and +24V.

[0119] FIG. 7 is a diagrammatic view of an exemplary driving system **300** for controlling the pixel electrode array **302** of a four-particle color electrophoretic medium in accordance with one or more embodiments. The driving system **300** may be in the form of an integrated circuit. The elements of the array **302** are arranged in the form of a matrix having a plurality of data lines and a plurality of gate lines. Each element of the matrix contains a TFT for controlling the electrode potential of a corresponding electrode, and each TFT is connected to one of the gate lines and one of the data lines.

[0120] The controller shown comprises a microcontroller **304** including control logic and switching logic. It receives input data from the input data lines **322**. The microcontroller has an output for each data line of the matrix, providing a data signal. A data signal line **306** connects each output to a data line of the matrix. The microcontroller **304** also has an output for each gate line of the matrix, providing a gate line selection signal. A gate signal line **308** connects each output to a gate line of the matrix. A data line driver **310** and a gate line driver **312** are arranged in each data and gate signal line, respectively. The figure shows the signals lines only for those data lines and gate lines shown in the figure. The gate line drivers may be integrated in a single integrated circuit. Similarly, the data line drivers may be integrated in a

single integrated circuit (as a source driver). The integrated circuit may include the complete driver assembly together with the microcontroller.

[0121] The data line drivers provide the signal levels corresponding to voltages selected to drive the pixels. The gate line drivers provide the signals for selecting the gate line of which the electrodes are to be actuated. A sequence of voltages of one of the data line drivers **210** is shown in FIG. 7. When there is large enough positive voltage on the gate line then there is low impedance between the data line and pixel, so the voltage on the data line is transferred to the pixel. When there is a negative voltage on the TFT gate then the TFT is high impedance and voltage is stored on the pixel capacitor and not affected by the voltage on the data line. The figure shows four columns labelled n to $n+3$ and five rows labelled n to $n+4$.

[0122] As illustrated in FIG. 7, line-at-a-time addressing is used, in which one gate line n is high while all the others are low. The signals on all of the data lines are then transferred to all of the pixels in row n . At the end of the line time gate line n signal goes low and the next gate line $n+1$ goes high, so that data for the next line is transferred to the TFT pixels in row $n+1$. This continues with all of the gate lines being scanned sequentially so the whole matrix is driven. This is the same method that is used in almost all active matrix LCDs, such as mobile phone screens, laptop screens, and LCD-TVs, whereby TFTs control the voltage maintained across the liquid crystal layer, and in active matrix EPDs.

Example

[0123] The benefits of the full-color EPD discussed above addressed by waveforms having four voltage levels have been confirmed experimentally.

[0124] The experiments were performed on an EPD module comprising an electrophoretic medium of the type described above in connection with in FIGS. 2A-2E, i.e., containing white, cyan, magenta, and yellow particles depicted by W^- , C^{+++} , M^{++} , and Y^+ , respectively. The electrophoretic fluid was contained in microcups of 8 micron thickness.

[0125] The experiments were conducted using waveforms having varied numbers of voltage levels listed in Table 3.

Table 3. Waveform Voltage Levels

	Voltage Levels
3-Level	-24V, 0V, 24V
4-Level	-24V, -10V, 0V, 24V
4-Level*	-24V, 0V, 10V, 24V

5-Level	-24V, -10V, 0V, 10V, 24V
7-Level	-24V, -17V, -10V, 0V, 10V, 17V, 24V

[0126] The waveforms were tuned using the computer modeling methods described above.

[0127] The EPD module was addressed by each of the different waveforms, and the results were recorded and shown in Tables 4 to 8.

Table 4. 3-Level Waveform Results

+/- 24V, 0V				
	L*	a*	b*	C*
Black	19.0	-0.2	-8.7	8.7
Blue	25.8	-1.4	-21.2	21.2
Red	35.0	21.2	6.3	22.1
Magenta	36.8	21.6	-7.2	22.8
Green	43.9	-16.7	10.8	19.9
Cyan	45.9	-14.8	-7.0	16.4
Yellow	58.0	-8.2	33.7	34.7
White	62.7	-5.2	2.7	5.8
Gamut volume				16304

Table 5. 4-Level Waveform Results

+/- 24V, -10V, 0V				
	L*	a*	b*	C*
Black	20.0	-0.7	-10.5	10.6
Blue	26.8	-0.5	-23.1	23.1
Red	37.9	22.6	4.5	23.1
Magenta	41.0	29.3	-13.7	32.3
Green	45.3	-15.8	7.9	17.7
Cyan	47.9	-13.0	-9.2	15.9
Yellow	58.6	-8.4	27.1	28.4
White	62.0	0.2	-1.7	1.7
Gamut volume				17761

Table 6. 4-Level* Waveform Results

+/- 24V, +10V, 0V				
	L*	a*	b*	C*
Black	18.9	0.0	-8.8	8.8
Blue	25.6	-1.8	-20.4	20.5
Red	35.5	22.3	9.1	24.1
Magenta	37.3	21.5	-7.1	22.6

Green	40.0	-23.5	13.2	27.0
Cyan	46.3	-15.6	-7.1	17.1
Yellow	56.6	-11.2	40.4	41.9
White	62.6	-5.2	2.7	5.8
Gamut volume				21978

Table 7. 5-Level Waveform Results

+/-24V, +/-10V, 0V				
	L*	a*	b*	c*
Black	18.9	0.1	-8.7	8.7
Blue	26.9	-0.9	-23.6	23.6
Red	36.6	22.7	10.0	24.8
Magenta	40.5	31.6	-14.0	34.6
Green	42.5	-19.4	13.8	23.8
Cyan	48.4	-13.4	-9.1	16.2
Yellow	57.0	-10.2	37.9	39.3
White	62.4	-2.5	-0.4	2.5
Gamut volume				25789

Table 8. 7-Level Waveform Results

+/- 24V, +/-17V, +/-10V, 0V				
	L*	a*	b*	c*
Black	19.2	-0.1	-9.3	9.3
Blue	25.6	-0.4	-23.0	23.0
Red	35.6	21.7	9.2	23.6
Magenta	40.0	30.1	-13.6	33.1
Green	41.4	-20.3	14.2	24.8
Cyan	48.6	-12.8	-9.0	15.6
Yellow	57.0	-10.1	38.1	39.4
White	62.1	0.1	-0.4	0.4
Gamut volume				24701

[0128] The results show that the “4-Level” driving scheme with two negative voltage levels, one positive voltage level, and a near zero voltage level (Table 5) provides only a slight improvement over the “3-Level” driving scheme (Table 4) in color gamut volume, whereas the “4-Level*” driving scheme with two positive voltage levels, one negative voltage level, and a near zero voltage level (Table 6) approaches the quality obtainable with the “5-Level” driving scheme (Table 7). There is only a negligible difference between “5-Level” driving scheme and the “7-Level” driving scheme (Table 8).

[0129] Table 9 shows the details of the colors obtained with the various driving voltage schemes.

Table 9. Combined Results

		Voltage Levels				
		3-Level	4-Level	4-Level*	5-Level	7-Level
C*	Black	8.7	10.6	8.8	8.7	9.3
	Blue	21.2	23.1	20.5	23.6	23.0
	Red	22.1	23.1	24.1	24.8	23.6
	Magenta	22.8	32.3	22.6	34.6	33.1
	Green	19.9	17.7	27.0	23.8	24.8
	Cyan	16.4	15.9	17.1	16.2	15.6
	Yellow	34.7	28.4	41.9	39.3	39.4
	White	5.82	1.75	5.84	2.54	0.40

[0130] When the 4-Level* driving scheme with two positive voltages, one negative voltage, and a near zero voltage is used, the magenta and blue are reduced in quality relative to 5-Level driving scheme, as expected. However, this quality reduction can be tolerated, as the 5-Level driving scheme magenta in particular is out of balance with respect to the other colors, particularly yellow and green. Thus, some sacrifice of magenta quality is tolerable in exchange for improved yellow and green quality. This tradeoff is highlighted in FIG. 9, which shows the color gamut available for the various driving voltage schemes.

[0131] Thus, four particle pigment EPDs addressed by waveforms having exactly four voltage levels provides excellent color quality, while reducing costs for the display by using 2-bit rather than 3-bit per pixel data.

[0132] Having thus described several aspects and embodiments of the technology of this application, it is to be appreciated that various alterations, modifications, and improvements will readily occur to those of ordinary skill in the art. Such alterations, modifications, and improvements are intended to be within the spirit and scope of the technology described in the application. For example, those of ordinary skill in the art will readily envision a variety of other means and/or structures for performing the function and/or obtaining the results and/or one or more of the advantages described herein, and each of such variations and/or modifications is deemed to be within the scope of the embodiments described herein. Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments described herein. It is, therefore, to be understood that the foregoing embodiments are presented by way of example

only and that, within the scope of the appended claims and equivalents thereto, inventive embodiments may be practiced otherwise than as specifically described. In addition, any combination of two or more features, systems, articles, materials, kits, and/or methods described herein, if such features, systems, articles, materials, kits, and/or methods are not mutually inconsistent, is included within the scope of the present disclosure.

CLAIMS

1. A color electrophoretic display, comprising:
 - a light-transmissive electrode at a viewing surface;
 - a back electrode; and
 - an electrophoretic medium disposed between the light-transmissive electrode and the back electrode comprising:
 - a non-polar fluid; and
 - a multi-pigment particle system comprising four types of charged electrophoretic pigment particles dispersed in the non-polar fluid, the four types of charged electrophoretic pigment particles comprising:
 - a first type of particle having a first optical property and a first charge polarity;
 - a second type of particle having a second optical property and a second charge polarity, wherein the second charge polarity is opposite to the first charge polarity;
 - a third type of particle having a third optical property and the second charge polarity; and
 - a fourth type of particle having a fourth optical property and the second charge polarity;
- wherein the first, second, third, and fourth optical properties are different from one another;
- wherein multi-pigment particle system is directly addressable with push-pull waveforms applied to the back electrode, having pulse voltage levels selected from a set of exactly four different voltage levels to render any of eight primary colors of red, green, blue, cyan, magenta, yellow, black, and white at each pixel of the electrophoretic medium while holding the voltage on the light-transmissive electrode constant, said voltage levels comprising a first positive voltage, a second positive voltage lower than the first positive voltage, a near zero voltage, and a negative voltage.

2. The color electrophoretic display of Claim 1, wherein the first positive voltage is 15 to 30V, the second positive voltage is 5 to 15V, and the negative voltage is -15 to -30V.

3. The color electrophoretic display of Claim 1, wherein the first positive voltage is 24V, the second positive voltage is 10V, and the negative voltage is -24V.

4. The color electrophoretic display of Claim 1, wherein the push-pull waveforms comprise aperiodic voltage sequences.

5. The color electrophoretic display of Claim 1, wherein the first type of particle is a light scattering particle, and the second, third, and fourth types of particles are light-absorbing particles.

6. The color electrophoretic display of Claim 1, wherein the first type of particle is white, and the second, third, and fourth types of particles are selected from cyan, magenta, and yellow.

7. The color electrophoretic display of Claim 1, wherein the first type of particle is white, the second type of particle is cyan, the third type of particle is magenta, and the fourth type of particle is yellow.

8. The color electrophoretic display of Claim 1, wherein the first charge polarity is negative, and the second charge polarity is positive.

9. The color electrophoretic display of Claim 1, wherein the multi-pigment particle system has exactly four types of charged electrophoretic particles.

10. The color electrophoretic display of Claim 1, wherein the non-polar fluid includes charge-control additives.

11. The electrophoretic display of Claim 1, wherein the electrophoretic medium is encapsulated in capsules or contained in sealed microcells.

12. The electrophoretic display of Claim 1, wherein the electrophoretic display is configured for incorporation in a book reader, a portable computer, a tablet computer, a monitor, a phone, a smart card, a sign, a watch, jewelry, a shelf label, a panel for a vehicle, or a flash drive.

13. The color electrophoretic display of Claim 1, wherein the back electrode comprises a segmented electrode or an active matrix backplane including an array of pixel electrodes.

14. The color electrophoretic display of Claim 13, wherein the active matrix backplane includes an array of thin film transistors coupled to the pixel electrodes.

15. The color electrophoretic display of Claim 1, wherein the push-pull waveforms are determined using a trained computer model.

16. A system, comprising:

a color electrophoretic display according to any of Claims 1-15, wherein the back electrode comprises an active matrix backplane including an array of pixel electrodes;

a source driver;

at least one power supply; and

a controller for transmitting two bits of data per pixel to the source driver for controlling the source driver to apply selected voltages from the at least one power supply to selected pixel electrodes.

17. A method, comprising:

providing a color electrophoretic display, comprising a light-transmissive electrode at a viewing surface, a back electrode, and an electrophoretic medium disposed between the light-transmissive electrode and the back electrode, said electrophoretic medium comprising a non-polar fluid and a multi-pigment particle system comprising four types of charged electrophoretic pigment particles dispersed in the non-polar fluid, the four types of charged electrophoretic pigment particles comprising: a first type of particle having a first optical property and a first charge polarity; a second type of particle having a second optical property and a second charge polarity, wherein the second charge polarity is opposite to the first charge polarity; a third type of particle having a third optical property and the second charge polarity; and a fourth type of particle having a fourth optical property and the second charge polarity; wherein the first, second, third, and fourth optical properties are different from one another; and

directly addressing the multi-pigment particle system with push-pull waveforms applied to the back electrode having voltage levels selected from a set of exactly four different voltage levels to render any of eight primary colors of red, green, blue, cyan, magenta, yellow, black, and white at each pixel of the electrophoretic medium while holding the voltage on the light-transmissive electrode constant, said four different voltage levels comprise a first positive voltage, a second positive voltage lower than the first positive voltage, a near zero voltage, and a negative voltage.

18. The method according to any of the preceding claims, wherein the first positive voltage is 15 to 30V, the second positive voltage is 5 to 15V, and the negative voltage is -15 to -30V.

19. The method of Claim 17, wherein the first positive voltage is 24V, the second positive voltage is 10V, and the negative voltage is -24V.

20. The method of Claim 17, wherein the push-pull waveforms comprise aperiodic voltage sequences.

21. The method of Claim 17, wherein the first type of particle is a light scattering particle, and wherein the second, third, and fourth types of particles are light-absorbing particles.

22. The method of Claim 17, wherein the first type of particle is white, and the second, third, and fourth types of particles are selected from cyan, magenta, and yellow.

23. The method of Claim 17, wherein the first type of particle is white, the second type of particle is cyan, the third type of particle is magenta, and the fourth type of particle is yellow.

24. The method of Claim 17, wherein the first charge polarity is negative, and the second charge polarity is positive.

25. The method of Claim 17, wherein the multi-pigment particle system has exactly four types of charged electrophoretic particles.

26. The method of Claim 17, wherein the non-polar fluid includes charge-control additives.

27. The method of Claim 17, wherein the electrophoretic medium is encapsulated in capsules or contained in sealed microcells.

28. The method of Claim 17, wherein the electrophoretic display is configured for incorporation in a book reader, a portable computer, a tablet computer, a monitor, a phone, a smart card, a sign, a watch, jewelry, a shelf label, a panel for a vehicle, or a flash drive.

29. The method of Claim 17, wherein the back electrode comprises a segmented electrode or an active matrix backplane including an array of pixel electrodes.

30. The method according to Claim 29, wherein the active matrix backplane includes an array of thin film transistors coupled to the pixel electrodes.

31. The method of Claim 17, further comprising determining the push-pull waveforms using a trained computer model.

[illegible]

FIG. 1
PRIOR ART

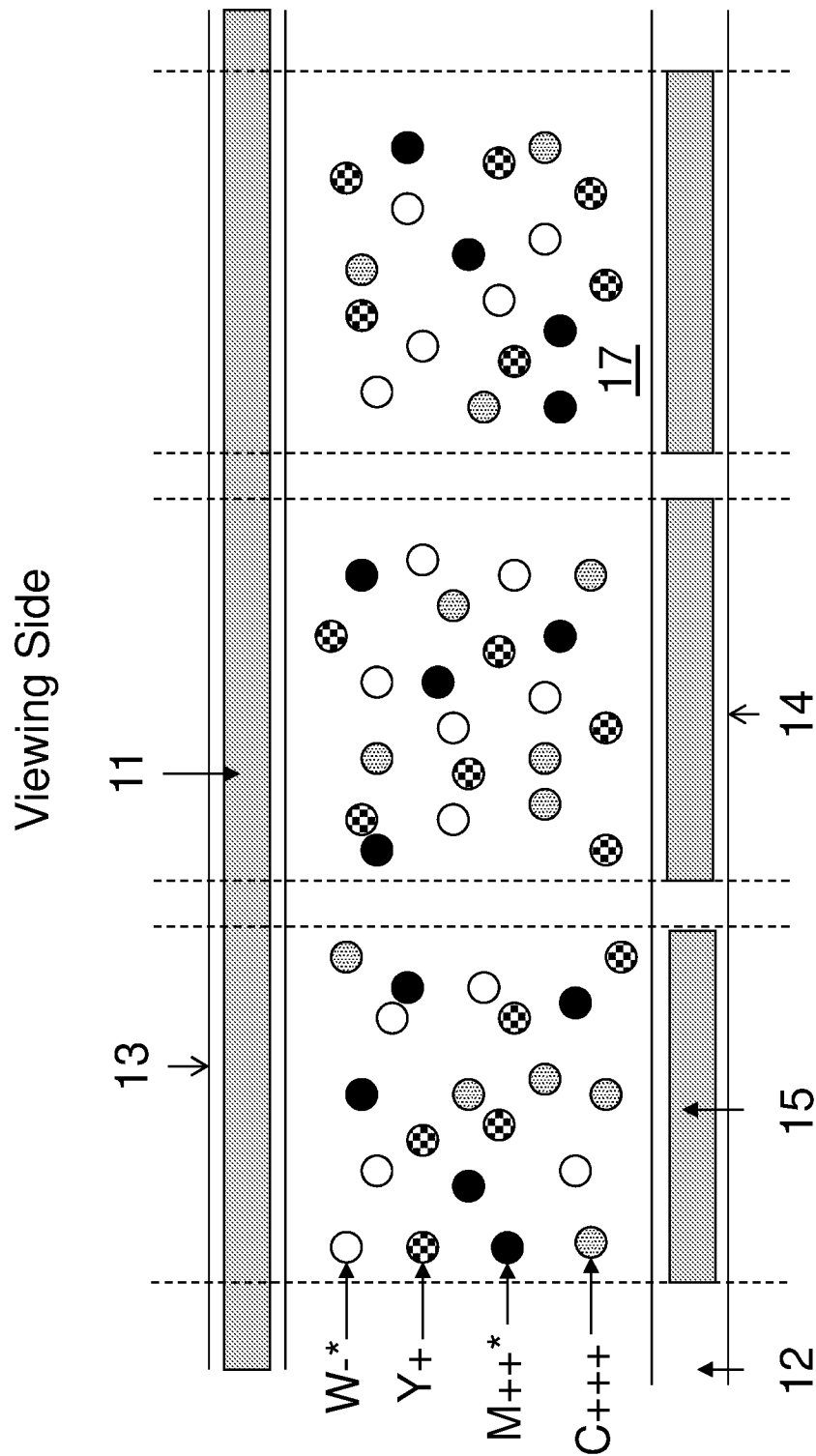


FIG. 2A
PRIOR ART

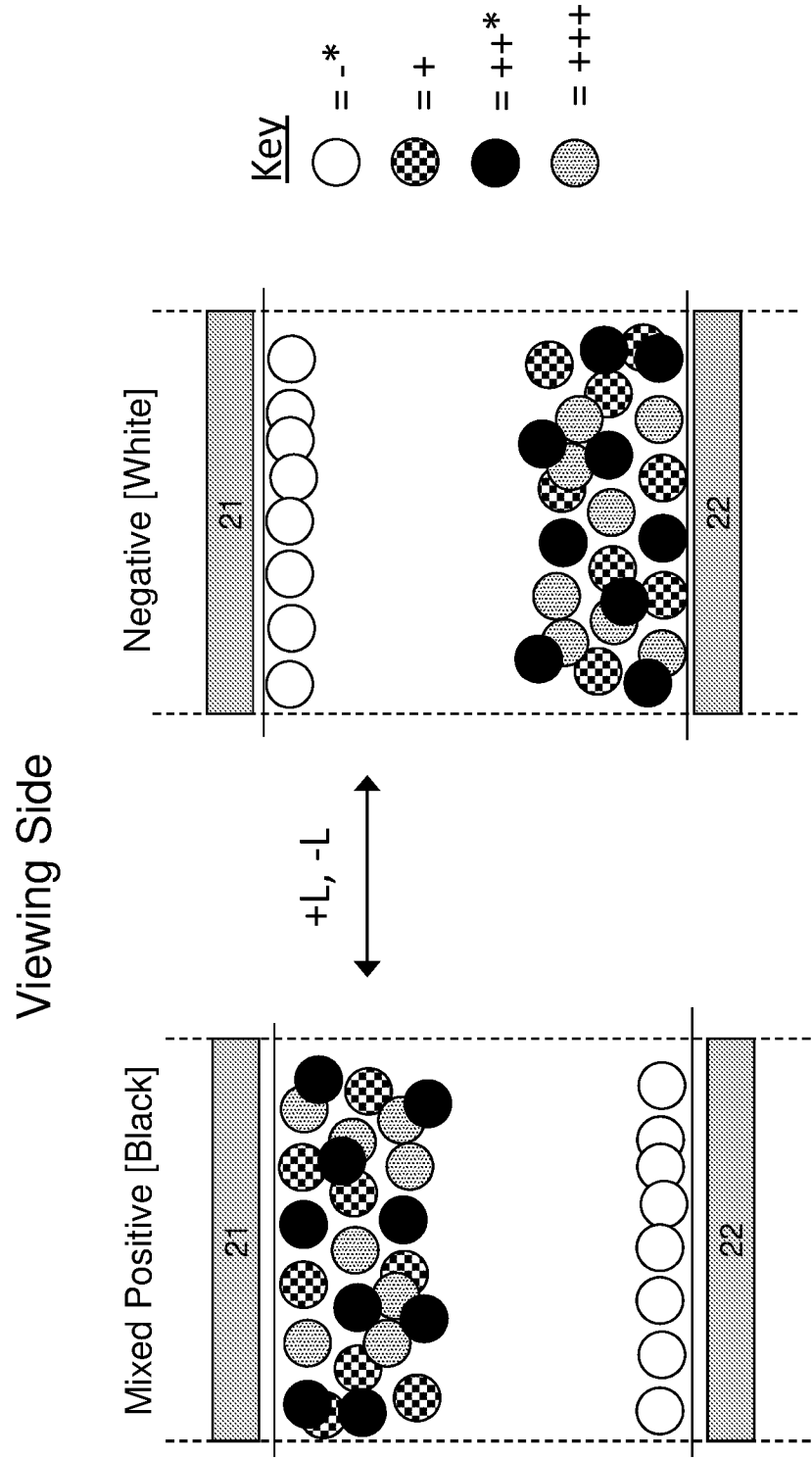


FIG. 2B
PRIOR ART

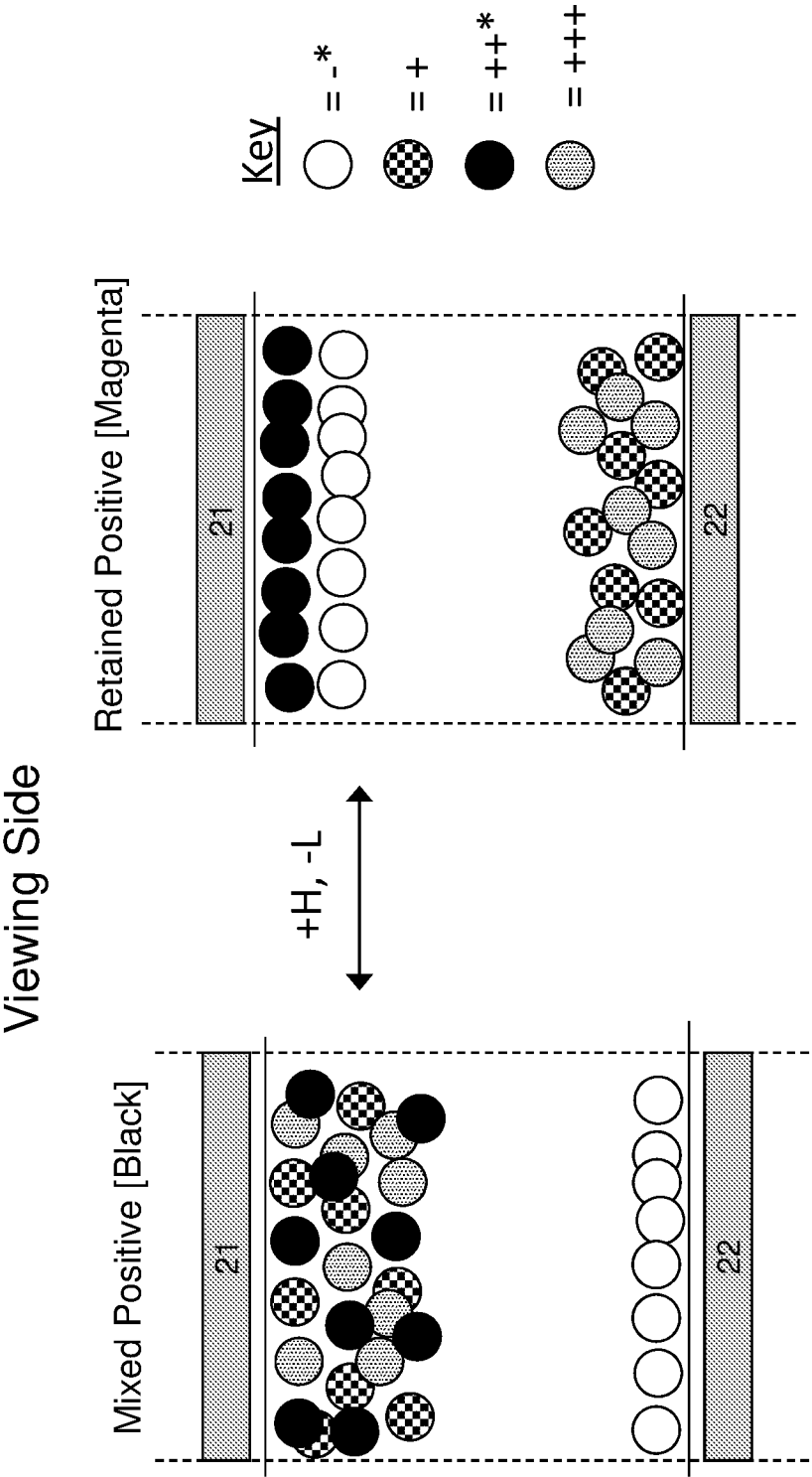


FIG. 2C
PRIOR ART

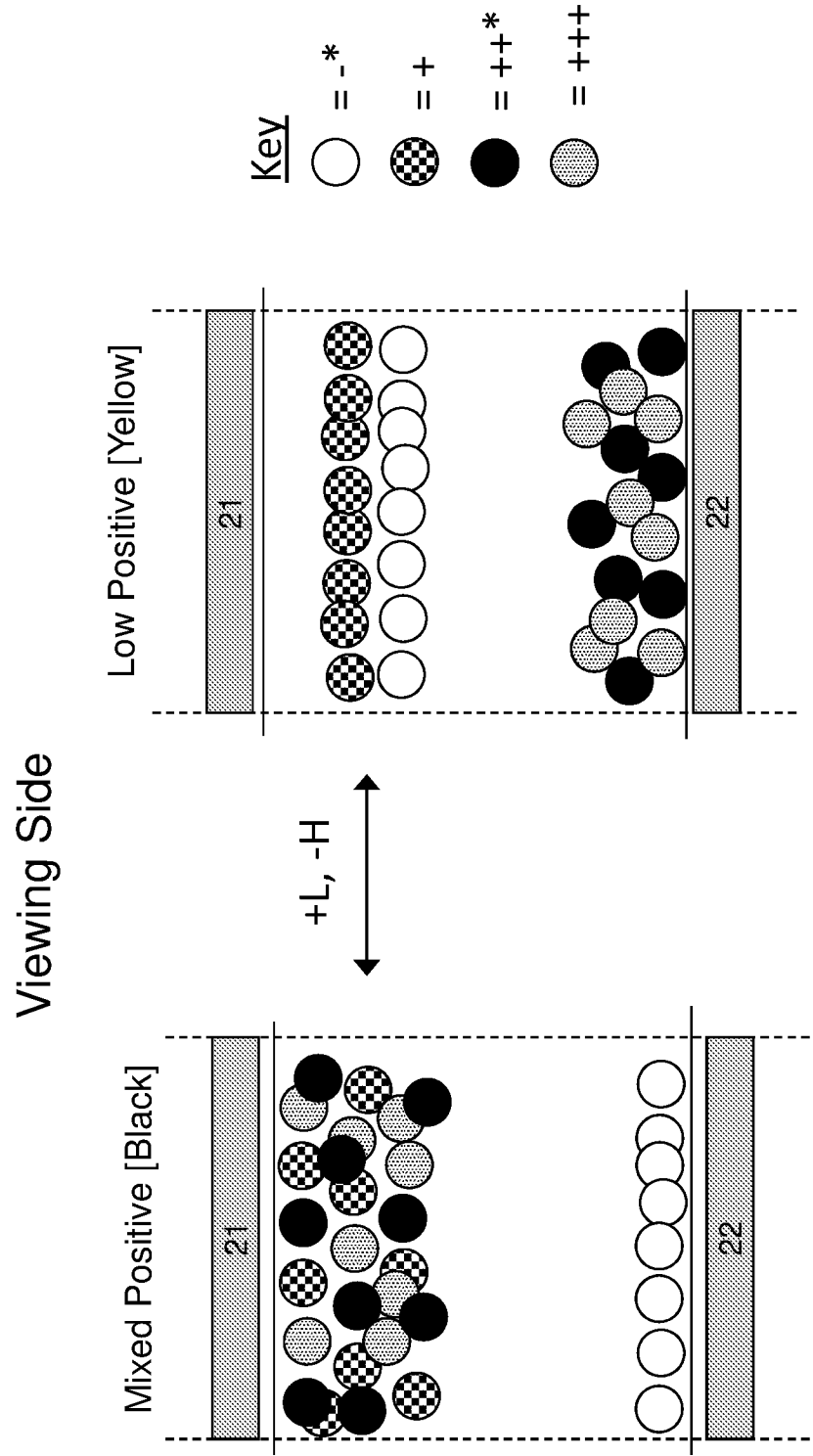


FIG. 2D
PRIOR ART

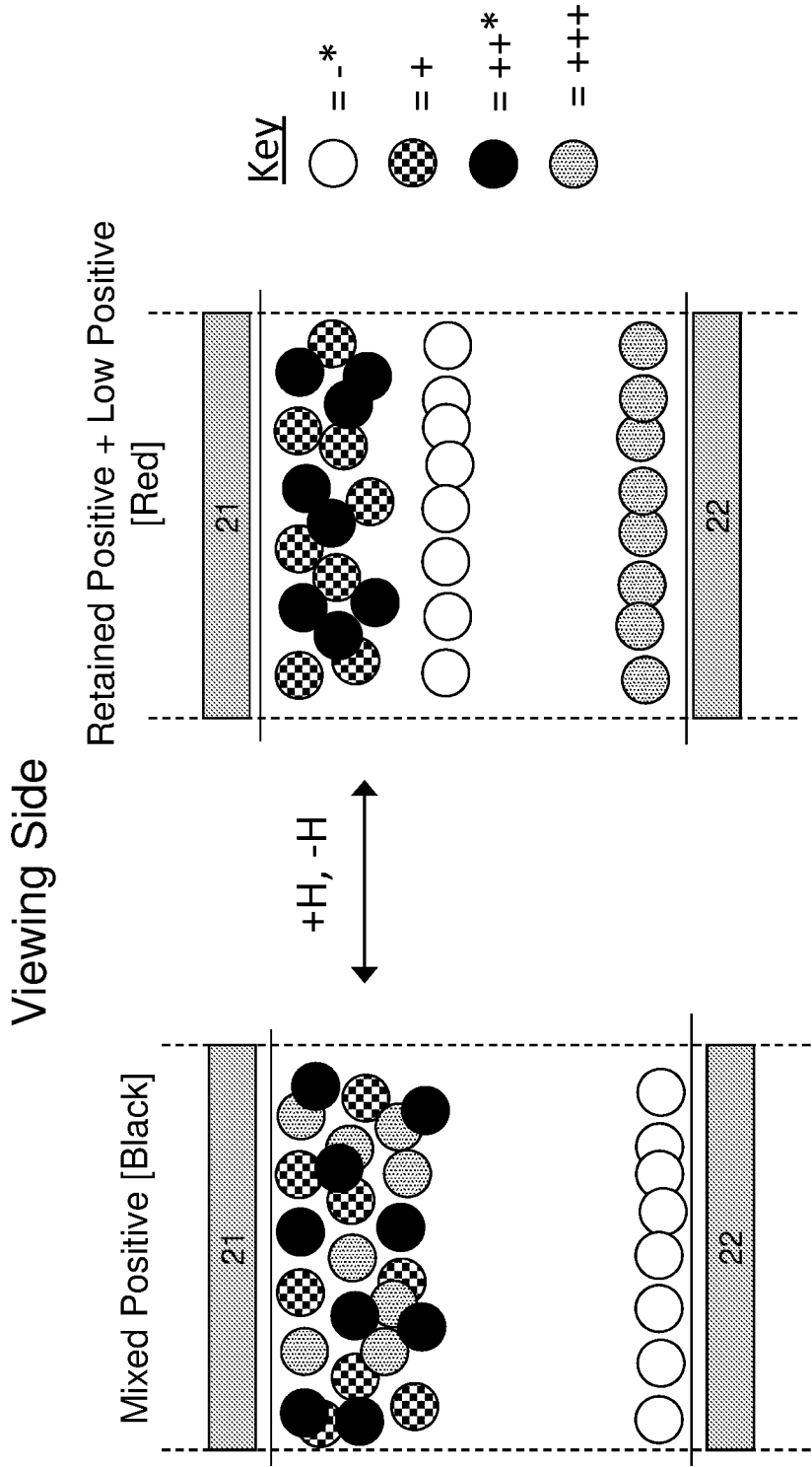


FIG. 2E
PRIOR ART

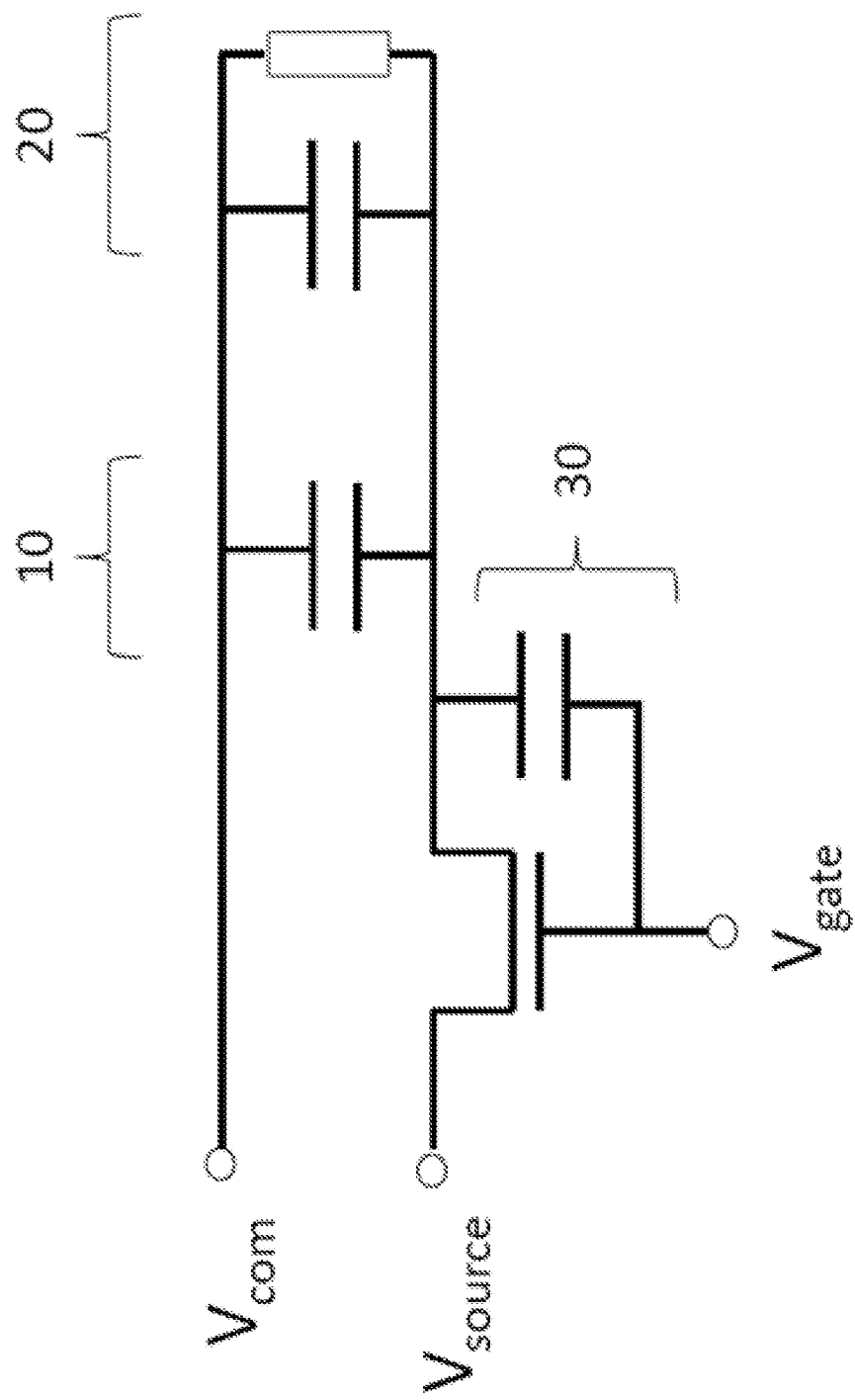


FIG. 3
PRIOR ART

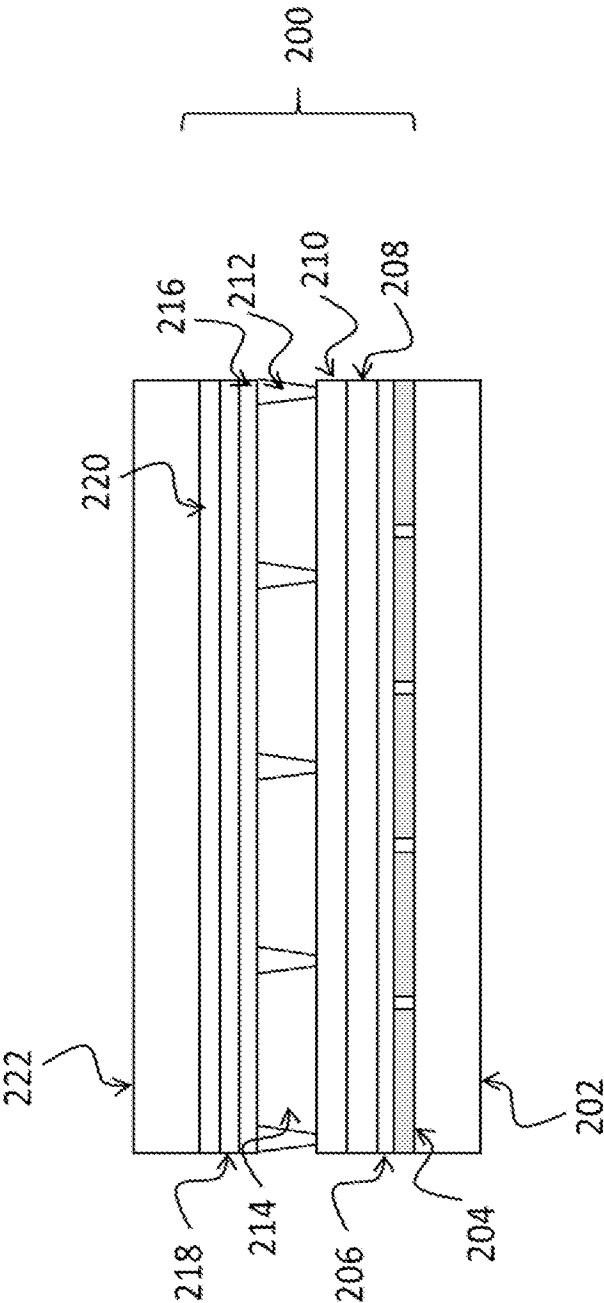


FIG. 4
PRIOR ART

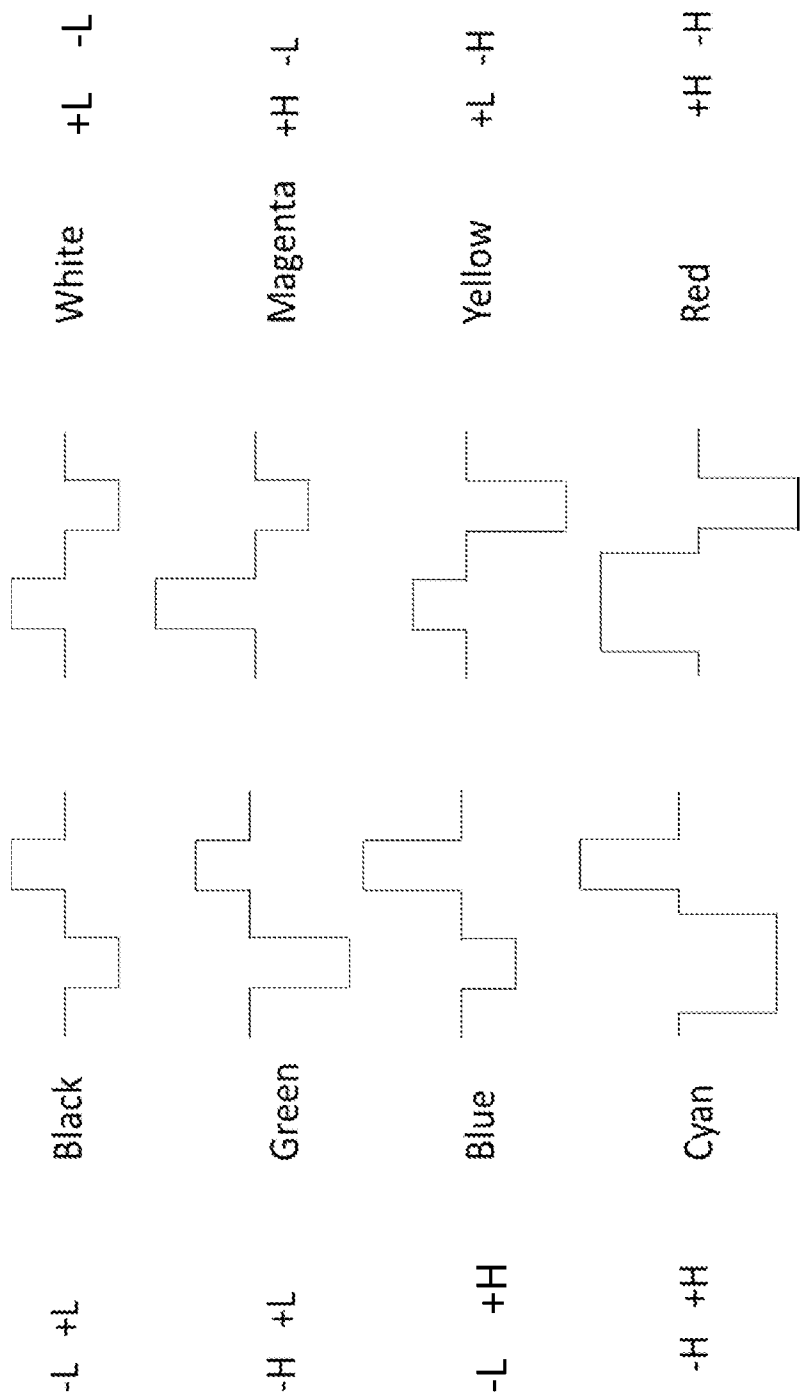
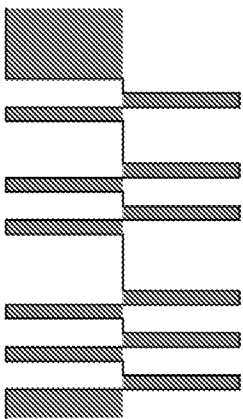
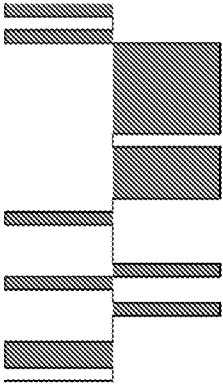


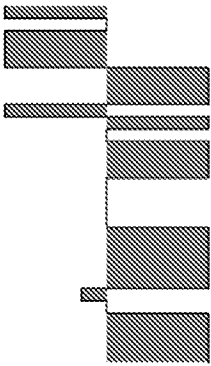
FIG. 5
PRIOR ART



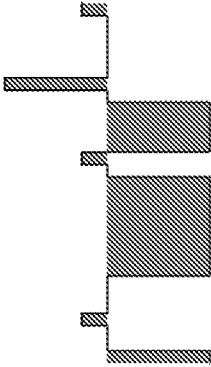
Black



Green



Blue

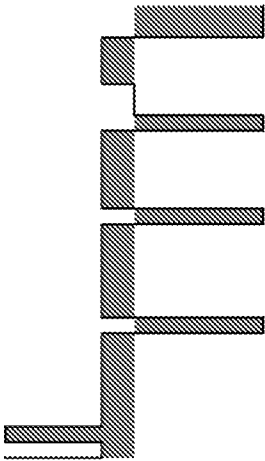


Cyan

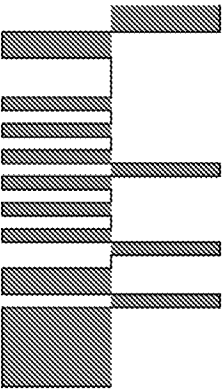
White



Yellow



Magenta



Red

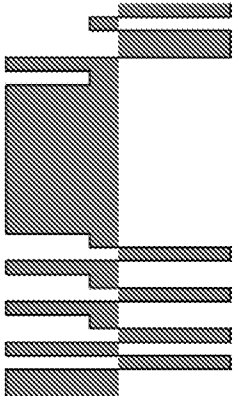


FIG. 6A

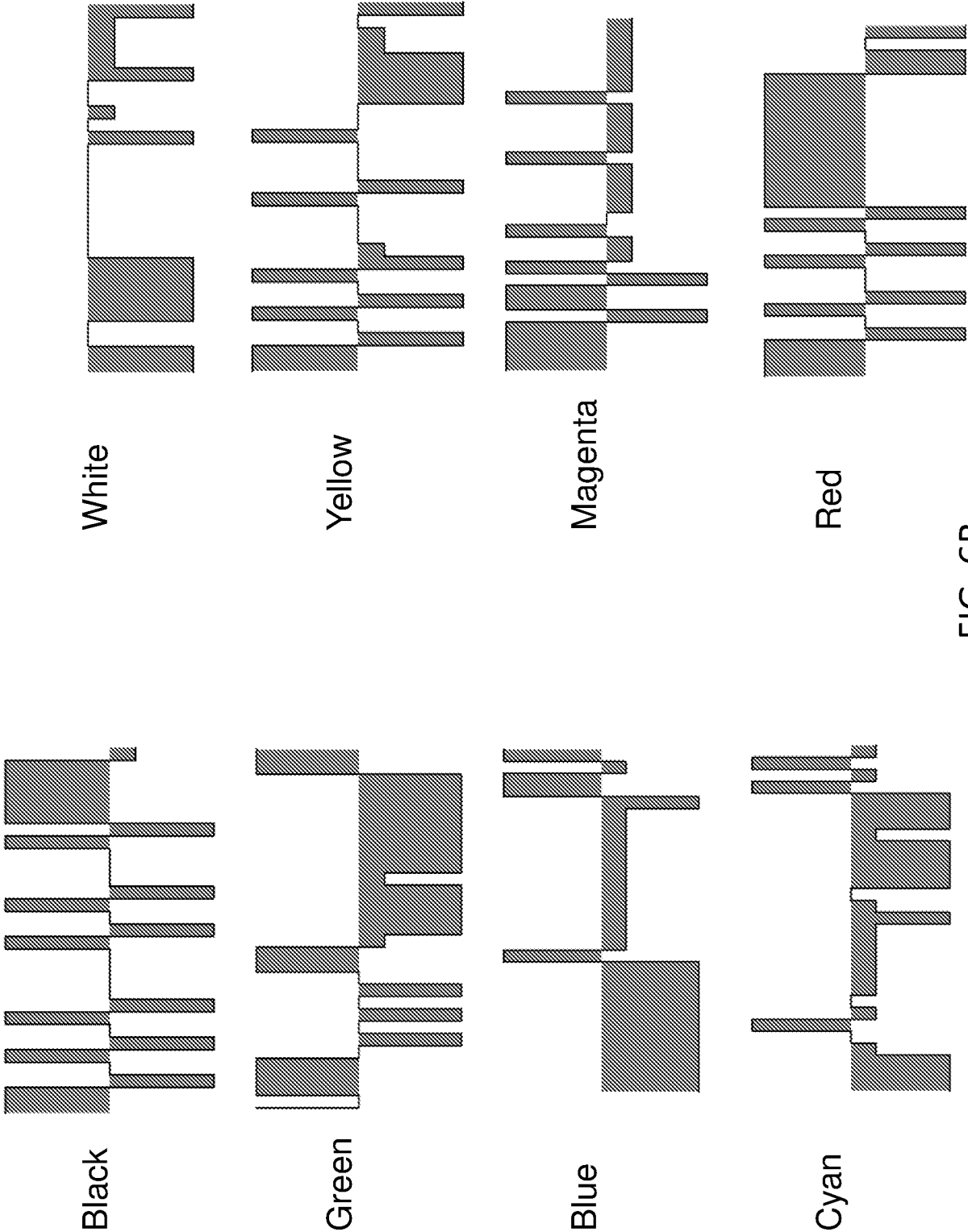
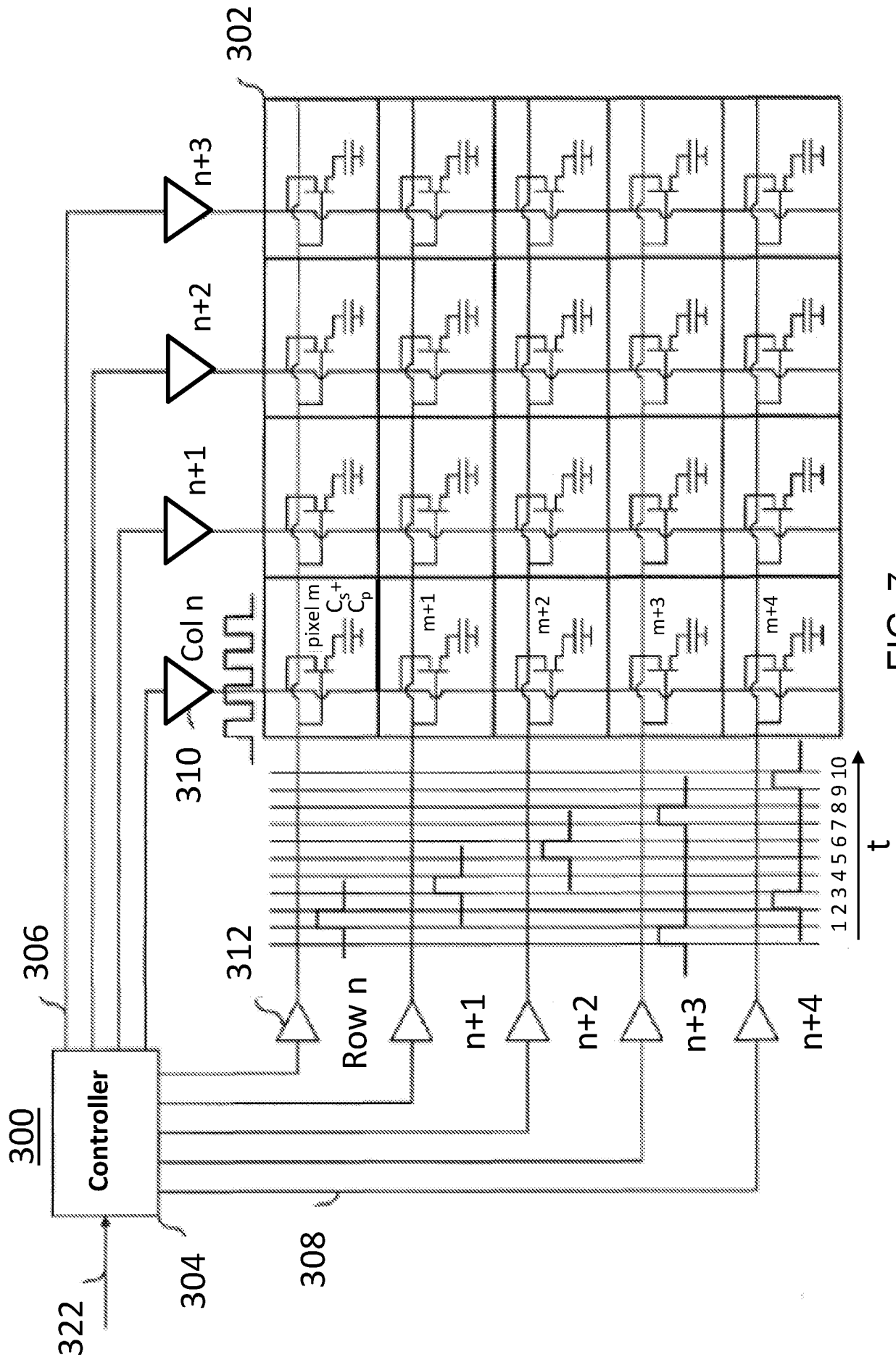


FIG. 6B



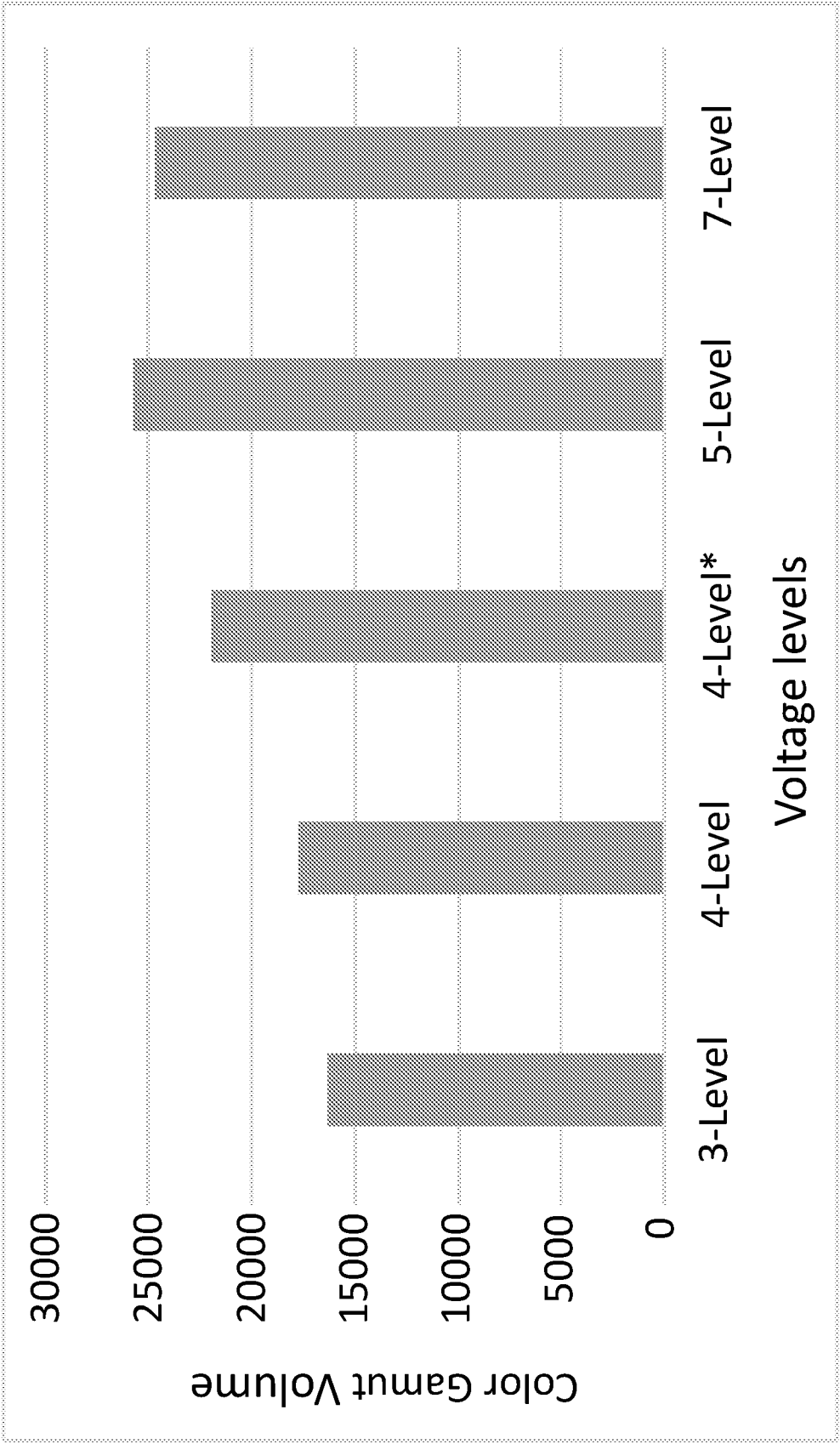


FIG. 8

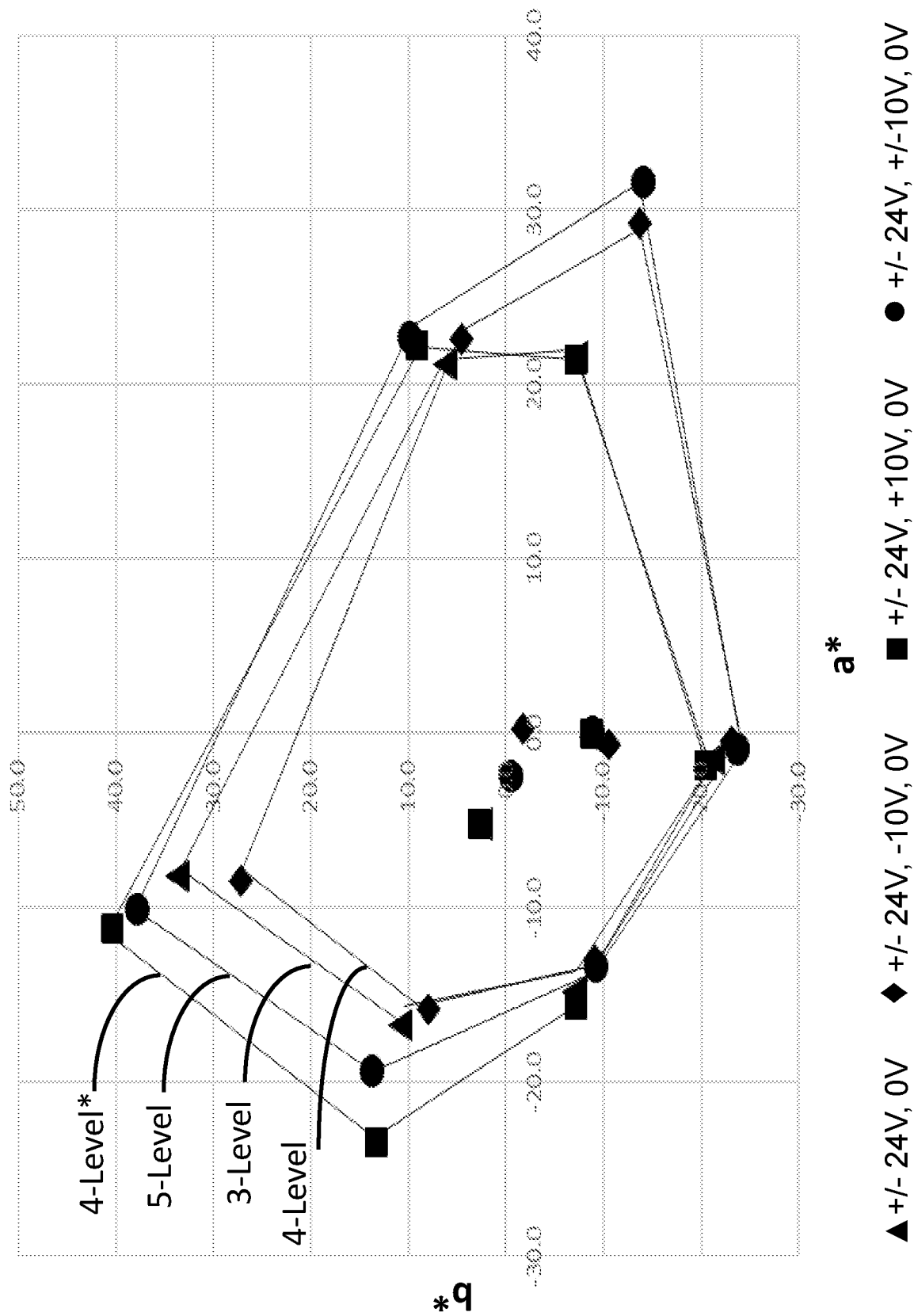


FIG. 9

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2024/031592

A. CLASSIFICATION OF SUBJECT MATTER INV. G09G3/34 G02F1/1685 G02F1/1675 ADD.		
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) G09G 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 practicable, 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.
A	US 2022/082896 A1 (TELFER STEPHEN J [US] ET AL) 17 March 2022 (2022-03-17) abstract; figures 1-5 paragraph [0049] - paragraph [0100] -----	1 - 31
A	US 2022/084473 A1 (TELFER STEPHEN J [US] ET AL) 17 March 2022 (2022-03-17) figures 1-6 paragraph [0048] - paragraph [0081] -----	1 - 31
A	US 2016/085132 A1 (TELFER STEPHEN J [US] ET AL) 24 March 2016 (2016-03-24) figures 1-6 paragraph [0040] - paragraph [0060] -----	1 - 31
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Date of the actual completion of the international search 23 September 2024		Date of mailing of the international search report 01/10/2024
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer Kentischer, Florian

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