

US00RE40787E

US RE40,787 E

(19) United States

(12) Reissued Patent

Martin et al.

(45) Date of Reissued Patent: Jun. 23, 2009

(54) MULTILAYER PLASTIC SUBSTRATES

- (75) Inventors: Peter M. Martin, Kennewick, WA (US);
 Gordon L. Graff, West Richland, WA (US); Mark E. Gross, Pasco, WA (US);
 Michael G. Hall, West Richland, WA (US); Eric S. Mast, Richland, WA (US)
- (73) Assignee: **Battelle Memorial Institute**, Columbus, OH (US)
- (21) Appl. No.: 10/889,640
- (22) Filed: Jul. 12, 2004

Related U.S. Patent Documents

Reissue of:

(64)	Patent No.:	6,623,861
	Issued:	Sep. 23, 2003
	Appl. No.:	09/835,768
	Filed:	Apr. 16, 2001

- U.S. Applications:
- (63) Continuation-in-part of application No. 09/427,138, filed on Oct. 25, 1999, now Pat. No. 6,522,067.
- (51) Int. Cl. *B32B 27/36* (2006.01)

(56) **References Cited**

U.S. PATENT DOCUMENTS

2,382,432 A	8/1945	McManus et al.
2,384,500 A	9/1945	Stoll
3,475,307 A	10/1969	Karl-Heinz et al.
3,607,365 A	9/1971	Lindlof
3,941,630 A	3/1976	Larrabee
4,061,835 A	12/1977	Poppe et al.

(Continued)

1<u>00</u>

FOREIGN PATENT DOCUMENTS

704297	2/1968
2353506 A	A1 5/2000
196 03 746 A	A1 4/1997
696 15 510 T	5/2002
0 147 696 E	31 7/1985
0 299 753 A	A2 1/1989
0 299 753	1/1989
0 340 935	11/1989
0 340 935 A	A2 11/1989
0 390 540 E	3/1990

(10) Patent Number:

BE

CA

DE

DE

EP

EP EP

EP EP

EP

(Continued)

OTHER PUBLICATIONS

Clark I. Bright, et al., Transparent Barrier Coatings Based on ITO for Flexible Plastic Displays, Oct. 17–19, 1999, pp. 247–264, Tucson, Arizona.

Akedo et al., "LP–5: Lake–News Poster: Plasma–CVD SiNx/Plasma–Polymerized CNx:H Multi–layer Passivation Films for Organic Light Emmitting Diods", SID 03 Digest.* Chwang et al., "Thin Film encapsulated flexible organic electroluminescent displays", American Institute of Physics, 2003.*

Notification of Transmittal of the International Search Report Or The Declaration, Mar. 3, 2000, PCT/US99/29853. Graupner, W. et al.; "High Resolution Color Organic Light Emitting Diode Microdisplay Fabrication Method", SPIE Proceedings 4207; 11–19 (2000); pp. 1–9.

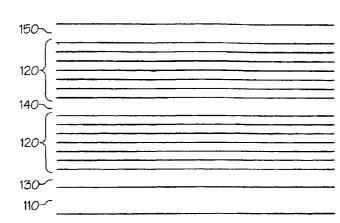
(Continued)

Primary Examiner—Milton I. Cano Assistant Examiner—Tamra L. Dicus (74) Attorney, Agent, or Firm—Dinsmore & Shohl LLP

(57) ABSTRACT

A multilayer plastic substrate. The substrate comprises a plurality of thin film layers of at least one polymer, the plurality of thin film layers being adjacent to one another and having sufficient strength to be self-supporting, wherein the multilayer plastic substrate has an average visible light transmittance of greater than about 80%.

23 Claims, 1 Drawing Sheet



U.S. PATENT DOCUMENTS

	0.8.	PATENT	DOCUMENTS
4,098,965	Α	7/1978	Kinsman
4,266,223	Α	5/1981	Frame
4,283,482	Α	8/1981	Hattori et al.
4,313,254	Α	2/1982	Feldman et al.
4,426,275	А	1/1984	Meckel et al.
4,521,458	A	6/1985	Nelson
4,537,814	A	8/1985	Itoh et al.
4,555,274	A	11/1985	Kitajima et al.
4,557,978	A A	12/1985 2/1986	Mason Christen
4,572,845 4,581,337	A	4/1986	Frey et al.
4,624,867	A	11/1986	lijima et al.
4,695,618	Ā	9/1987	Mowrer
4,710,426	Α	12/1987	Stephens
4,722,515	Α	2/1988	Ham
4,768,666	Α	9/1988	Kessler
4,842,893	Α	6/1989	Yializis et al.
4,843,036	A	6/1989	Schmidt et al.
4,854,186	A	8/1989	Grolig et al.
4,855,186	A	8/1989	Grolig et at.
4,889,609	A A	12/1989	Cannella Harada et al.
4,913,090 4,931,158	A	4/1990 6/1990	Bunshah et al.
4,934,315	A	6/1990	Linnebach et al.
4,954,371	A	9/1990	Yializis
4,977,013	Α	12/1990	Ritchie et al.
5,032,461	Α	7/1991	Shaw et al.
5,036,249	Α	7/1991	Pike-Biegunski et al.
5,047,131	Α	9/1991	Wolfe et al.
5,059,861	Α	10/1991	Littman et al.
5,124,204	A	6/1992	Yamashita et al.
5,189,405	A	2/1993	Yamashita et al.
5,203,898	A A	4/1993	Carpenter et al. Kirlin et al.
5,204,314 5,237,439	A	4/1993 8/1993	Misono et al.
5,260,095	Ā	11/1993	Affinito
5,336,324	Ā	8/1994	Stall et al.
5,354,497	Α	10/1994	Fukuchi et al.
5,356,947	Α	10/1994	Ali et al.
5,376,467	Α	12/1994	Abe et al.
5,393,607	Α	2/1995	Kawasaki et al.
5,395,644	A	3/1995	Affinito
5,402,314	A A	3/1995	Amago et al.
5,427,638 5,440,446	A	6/1995 8/1995	Goetz et al. Shaw et al.
5,451,449		9/1995	Shetty et al.
5,461,545	A	10/1995	Leroy et al.
5,464,667	A	11/1995	Kohler et al.
5,510,173	Α	4/1996	Pass et al.
5,512,320	Α	4/1996	Turner et al.
5,536,323	Α	7/1996	Kirlin et al.
5,547,508	A	8/1996	Affinito
5,554,220	A	9/1996	Forrest et al.
5,576,101 5,578,141	A A	11/1996	Saitoh et al.
5,607,789	A	11/1996 3/1997	Mori et al. Treger et al.
5,620,524	A	4/1997	Fan et al.
5,629,389	Ā	5/1997	Roitman et al.
5,652,192	A	7/1997	Matson et al.
5,654,084	Α	8/1997	Egert
5,660,961	Α	8/1997	Yu
5,665,280	Α	9/1997	Tropsha
5,681,615	A	10/1997	Affinito et al.
5,681,666	A	10/1997	Treger et al.
5,684,084	A	11/1997	Lewin et al.
5,686,360 5,693,956	A A	11/1997 12/1997	Harvey, III et al. Shi et al.
5,695,564	A	12/1997	Imahashi
5,711,816	A	1/1998	Kirlin et al.
5,725,909	A	3/1998	Shaw et al.

5,731,661 A	3/1998	So et al.
5,736,207 A	4/1998	Walther et al.
5,747,182 A	5/1998	Friend et al.
5,757,126 A	5/1998	Harvey, et al.
5,759,329 A	6/1998	Krause et al.
5,771,177 A	6/1998	Tada et al.
5,771,562 A	6/1998	Harvey, III et al.
5,782,355 A	7/1998	Katagiri et al.
5,792,550 A	8/1998	Phillips et al.
5,795,399 A	8/1998	Hasegawa et al.
5,811,177 A	9/1998	Shi et al.
5,811,183 A	9/1998	Shaw et al.
5,821,138 A	10/1998	Yamazaki et al.
5,821,692 A	10/1998	Rogers et al.
5,844,363 A	12/1998	Gu et al.
5,869,791 A	2/1999	Young
5,872,355 A	2/1999	Hueschen
5,891,554 A	4/1999	Hosokawa et al.
5,895,228 A	4/1999	Biebuyck et al.
5,902,641 A	5/1999	Affinito et al.
5,902,688 A	5/1999	Antoniadis et al.
5,904,958 A	5/1999	Dick et al.
· · ·		Yializis et al.
· · ·	6/1999	
5,919,328 A	7/1999	Tropsha et al.
5,920,080 A	7/1999	Jones
5,922,161 A	7/1999	Wu et al.
5,929,562 A	7/1999	Pichler
5,934,856 A	8/1999	Asakawa et al.
5,945,174 A	8/1999	Shaw et al.
5,948,552 A	9/1999	Antoniadis et al.
5,952,778 A	9/1999	Haskal et al.
5,955,161 A	9/1999	Tropsha
5,965,907 A	10/1999	Huang et al.
5,968,620 A	10/1999	Harvey et al.
5,994,174 A	11/1999	Carey et al.
5,996,498 A	12/1999	Lewis
6,013,337 A	1/2000	Knors
6,040,017 A	3/2000	Mikhael et al.
6,045,864 A	4/2000	Lyons et al.
6,066,826 A	5/2000	Yializis
6,083,313 A	7/2000	Venkatraman et al.
6,083,628 A	7/2000	Yializis
6,084,702 A	7/2000	Byker et al.
6,087,007 A	7/2000	Fujii et al
6,092,269 A	7/2000	Yializis et al.
6,106,627 A	8/2000	Yializis
6,117,266 A	9/2000	Horzel et al.
6,118,218 A	9/2000	Yializis et al.
6,146,225 A	11/2000	Sheats et al.
6,146,462 A	11/2000	Yializis et al.
6,150,187 A	11/2000	Zyung et al.
6,165,566 A	12/2000	Tropsha
6,178,082 B1	1/2001	Farooq et al.
6,195,142 B1	2/2001	Gyotoku et al.
6,198,217 B1	3/2001	Suzuki et al.
6,198,220 B1	3/2001	Jones et al.
6,203,898 B1	3/2001	Kohler et al.
6,207,238 B1	3/2001	Affinito
6,207,239 B1	3/2001	Affinito
6,214,422 B1	4/2001	Yializis
6,217,947 B1	4/2001	Affinito
6,224,948 B1	5/2001	Affinito
6,228,434 B1	5/2001	Affinito
6,228,436 B1	5/2001	Affinito
6,231,939 B1	5/2001	Shaw et al.
6,264,747 B1	7/2001	Shaw et al.
6,268,695 B1	7/2001	Affinito
6,274,204 B1	8/2001	Affinito
6,322,860 B1	11/2001	Stein et al.
6,333,065 B1	12/2001	Arai et al.
6,348,237 B2	2/2002	Kohler et al.

6,350,034 B1		
	2/2002	Fleming et al.
6,352,777 B1	3/2002	Bulovic et al.
6,358,570 B1	3/2002	Affinito
6,361,885 B1	3/2002	Chou
6,397,776 B1	6/2002	Yang et al.
6,413,645 B1	7/2002	Graff et al.
6,416,872 B1	7/2002	Maschwitz
6,420,003 B2	7/2002	Shaw et al.
6,436,544 B1	8/2002	Veyrat et al.
6,460,369 B2	10/2002	Hosokawa
6,465,953 B1	10/2002	Duggal
6,468,595 B1	10/2002	Mikhael et al.
6,469,437 B1	10/2002	Parthasarathy et al.
6,492,026 B1 6,497,598 B2	12/2002 12/2002	Graff et al. Affinito
6,497,598 B2 6,497,924 B2	12/2002	Affinito et al.
6,509,065 B2	1/2002	Affinito
6,512,561 B1	1/2003	Terashiat et al.
6,522,067 B1	2/2003	Graff et al.
6,537,688 B2	3/2003	Silvernail et al.
6,544,600 B2	4/2003	Affinito et al.
6,548,912 B1	4/2003	Graff et al.
6,569,515 B2	5/2003	Hebrink et al.
6,570,325 B2	5/2003	Graff et al.
6,573,652 B1	6/2003 6/2003	Graff et al.
6,576,351 B2 6,592,969 B1	7/2003	Silvernail Burroughes et al
6,597,111 B2	7/2003	Burroughes et al. Silvernail et al.
6,613,395 B2	9/2003	Affinito et al.
6,614,057 B2	9/2003	Silvernail et al.
6,624,568 B2	9/2003	Silvernail et al.
6,627,267 B2	9/2003	Affinito
6,628,071 B1	9/2003	Su
6,653,780 B2	11/2003	Sugimoto et al.
6,656,537 B2	12/2003	Affinito et al.
6,660,409 B1	12/2003	Komatsu et al.
6,664,137 B2 6,681,716 B2	12/2003 1/2004	Weaver Schaepkens
6,681,716 B2 6,720,203 B2	4/2004	Carcia et al.
6,734,625 B2	5/2004	Vong et al.
6,737,753 B2	5/2004	Kumar et al.
6,743,524 B2	6/2004	Schaepkens
6,749,940 B1	6/2004	Terasaki et al.
6,765,351 B2	7/2004	Forrest et al.
6,803,245 B2	10/2004	Auch et al.
6,811,829 B2		
	11/2004	Affinito et al.
6,815,887 B2	11/2004	Affinito et al. Lee et al.
6,815,887 B2 6,818,291 B2	11/2004 11/2004	Affinito et al. Lee et al. Funkenbusch et al.
6,815,887 B2 6,818,291 B2 6,835,950 B2	11/2004 11/2004 12/2004	Affinito et al. Lee et al. Funkenbusch et al. Brown et al.
6,815,887 B2 6,818,291 B2 6,835,950 B2 6,836,070 B2	11/2004 11/2004 12/2004 12/2004	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al.
6,815,887 B2 6,818,291 B2 6,835,950 B2 6,836,070 B2 6,837,950 B1	11/2004 11/2004 12/2004 12/2004 1/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard
6,815,887 B2 6,818,291 B2 6,835,950 B2 6,836,070 B2 6,837,950 B1 6,864,629 B2	11/2004 11/2004 12/2004 12/2004	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al.
6,815,887 B2 6,818,291 B2 6,835,950 B2 6,836,070 B2 6,837,950 B1	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard
6,815,887 B2 6,818,291 B2 6,835,950 B2 6,836,070 B2 6,837,950 B1 6,864,629 B2 6,866,901 B2 6,866,901 B2 6,867,539 B1 6,872,114 B2	11/2004 11/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al.
6,815,887 B2 6,818,291 B2 6,835,950 B2 6,836,070 B2 6,837,950 B1 6,864,629 B2 6,866,901 B2 6,867,539 B1 6,872,114 B2 6,872,248 B2	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Mizutani et al.
6,815,887 B2 6,818,291 B2 6,835,950 B2 6,836,070 B2 6,837,950 B1 6,864,629 B2 6,866,901 B2 6,867,539 B1 6,872,114 B2 6,872,248 B2 6,872,248 B2	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Mizutani et al. Yang et al.
$\begin{array}{c} 6,815,887 & B2\\ 6,818,291 & B2\\ 6,835,950 & B2\\ 6,836,070 & B2\\ 6,837,950 & B1\\ 6,864,629 & B2\\ 6,866,901 & B2\\ 6,867,539 & B1\\ 6,872,114 & B2\\ 6,872,248 & B2\\ 6,872,248 & B2\\ 6,872,428 & B2\\ 6,878,467 & B2\\ \end{array}$	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005 3/2005 3/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Yang et al. Chung et al.
$\begin{array}{c} 6,815,887 & B2\\ 6,818,291 & B2\\ 6,835,950 & B2\\ 6,835,950 & B2\\ 6,837,950 & B1\\ 6,864,629 & B2\\ 6,866,901 & B2\\ 6,867,539 & B1\\ 6,872,114 & B2\\ 6,872,248 & B2\\ 6,872,428 & B2\\ 6,872,428 & B2\\ 6,878,467 & B2\\ 6,888,305 & B2\\ \end{array}$	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005 3/2005 5/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Mizutani et al. Yang et al. Chung et al. Weaver
$\begin{array}{c} 6,815,887 & B2\\ 6,818,291 & B2\\ 6,835,950 & B2\\ 6,836,070 & B2\\ 6,837,950 & B1\\ 6,864,629 & B2\\ 6,866,901 & B2\\ 6,867,539 & B1\\ 6,872,114 & B2\\ 6,872,248 & B2\\ 6,872,428 & B2\\ 6,872,428 & B2\\ 6,878,467 & B2\\ 6,888,305 & B2\\ 6,888,307 & B2\\ \end{array}$	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005 5/2005 5/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Yang et al. Chung et al. Weaver Silvernail et al.
$\begin{array}{ccccccccc} 6,815,887 & B2\\ 6,818,291 & B2\\ 6,835,950 & B2\\ 6,836,070 & B2\\ 6,837,950 & B1\\ 6,864,629 & B2\\ 6,866,901 & B2\\ 6,867,539 & B1\\ 6,872,114 & B2\\ 6,872,248 & B2\\ 6,872,428 & B2\\ 6,878,467 & B2\\ 6,888,305 & B2\\ 6,888,307 & B2\\ 6,891,330 & B2\\ \end{array}$	11/2004 11/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005 3/2005 5/2005 5/2005 5/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Yang et al. Chung et al. Chung et al. Weaver Silvernail et al. Duggal et al.
$\begin{array}{ccccccc} 6,815,887 & B2\\ 6,818,291 & B2\\ 6,835,950 & B2\\ 6,837,950 & B1\\ 6,864,629 & B2\\ 6,866,901 & B2\\ 6,867,539 & B1\\ 6,872,114 & B2\\ 6,872,248 & B2\\ 6,872,248 & B2\\ 6,872,428 & B2\\ 6,872,448 & B2\\ 6,878,467 & B2\\ 6,888,305 & B2\\ 6,888,307 & B2\\ 6,891,330 & B2\\ 6,897,474 & B2\\ \end{array}$	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005 5/2005 5/2005 5/2005 5/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Yang et al. Chung et al. Weaver Silvernail et al. Duggal et al. Brown et al.
$\begin{array}{c} 6,815,887 & B2\\ 6,818,291 & B2\\ 6,835,950 & B2\\ 6,836,070 & B2\\ 6,837,950 & B1\\ 6,864,629 & B2\\ 6,866,901 & B2\\ 6,867,539 & B1\\ 6,872,114 & B2\\ 6,872,114 & B2\\ 6,872,248 & B2\\ 6,872,248 & B2\\ 6,872,428 & B2\\ 6,872,428 & B2\\ 6,878,467 & B2\\ 6,888,305 & B2\\ 6,888,307 & B2\\ 6,897,474 & B2\\ 6,897,474 & B2\\ 6,897,607 & B2\\ \end{array}$	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005 5/2005 5/2005 5/2005 5/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Yang et al. Chung et al. Chung et al. Weaver Silvernail et al. Duggal et al.
$\begin{array}{ccccccc} 6,815,887 & B2\\ 6,818,291 & B2\\ 6,835,950 & B2\\ 6,837,950 & B1\\ 6,864,629 & B2\\ 6,866,901 & B2\\ 6,867,539 & B1\\ 6,872,114 & B2\\ 6,872,248 & B2\\ 6,872,248 & B2\\ 6,872,428 & B2\\ 6,872,448 & B2\\ 6,878,467 & B2\\ 6,888,305 & B2\\ 6,888,307 & B2\\ 6,891,330 & B2\\ 6,897,474 & B2\\ \end{array}$	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005 5/2005 5/2005 5/2005 5/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Yang et al. Chung et al. Yang et al. Chung et al. Weaver Silvernail et al. Duggal et al. Brown et al.
$\begin{array}{ccccccc} 6,815,887 & B2\\ 6,818,291 & B2\\ 6,835,950 & B2\\ 6,836,070 & B2\\ 6,837,950 & B1\\ 6,864,629 & B2\\ 6,866,901 & B2\\ 6,867,539 & B1\\ 6,872,114 & B2\\ 6,872,114 & B2\\ 6,872,248 & B2\\ 6,872,248 & B2\\ 6,872,428 & B2\\ 6,872,428 & B2\\ 6,872,428 & B2\\ 6,873,467 & B2\\ 6,883,005 & B2\\ 6,888,307 & B2\\ 6,891,330 & B2\\ 6,897,474 & B2\\ 6,897,607 & B2\\ 6,905,769 & B2\\ \end{array}$	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 5/2005 5/2005 5/2005 5/2005 5/2005 5/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Mizutani et al. Yang et al. Chung et al. Weaver Silvernail et al. Duggal et al. Brown et al. Sugimoto et al. Komada
$\begin{array}{cccccccc} 6,815,887 & B2\\ 6,818,291 & B2\\ 6,835,950 & B2\\ 6,836,070 & B2\\ 6,837,950 & B1\\ 6,864,629 & B2\\ 6,866,901 & B2\\ 6,867,539 & B1\\ 6,872,114 & B2\\ 6,872,248 & B2\\ 6,872,248 & B2\\ 6,872,428 & B2\\ 6,872,428 & B2\\ 6,872,428 & B2\\ 6,878,467 & B2\\ 6,883,05 & B2\\ 6,888,307 & B2\\ 6,888,307 & B2\\ 6,891,330 & B2\\ 6,897,474 & B2\\ 6,897,474 & B2\\ 6,905,769 & B2\\ 6,923,702 & B2\\ \end{array}$	11/2004 11/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005 5/2005 5/2005 5/2005 5/2005 5/2005 5/2005 5/2005 5/2005 5/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Mizutani et al. Yang et al. Chung et al. Weaver Silvernail et al. Duggal et al. Brown et al. Sugimoto et al. Komada Graff et al.
6,815,887 B2 6,818,291 B2 6,835,950 B2 6,836,070 B2 6,837,950 B1 6,864,629 B2 6,867,539 B1 6,872,114 B2 6,872,248 B2 6,872,428 B2 6,897,474 B2 6,897,474 B2 6,997,607 B2 6,936,131 B2 6,975,067 B2 6,994,933 B1	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005 5/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Mizutani et al. Yang et al. Chung et al. Weaver Silvernail et al. Duggal et al. Brown et al. Sugimoto et al. Komada Graff et al. McCormick et al. McCormick et al. Bates
6,815,887 B2 6,818,291 B2 6,835,950 B2 6,837,950 B1 6,864,629 B2 6,867,539 B1 6,867,539 B1 6,872,114 B2 6,872,148 B2 6,872,428 B2 6,872,428 B2 6,872,428 B2 6,872,428 B2 6,872,428 B2 6,872,428 B2 6,872,428 B2 6,872,428 B2 6,873,467 B2 6,897,474 B2 6,897,607 B2 6,905,769 B2 6,923,702 B2 6,936,131 B2 6,975,067 B2 6,998,648 B2	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 5/2005 5/2005 5/2005 5/2005 5/2005 5/2005 5/2005 8/2005 8/2005 8/2005 2/2006 2/2006	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Mizutani et al. Yang et al. Chung et al. Weaver Silvernail et al. Duggal et al. Brown et al. Sugimoto et al. Komada Graff et al. McCormick et al. Bates Silvernail
6,815,887 B2 6,818,291 B2 6,835,950 B2 6,836,070 B2 6,837,950 B1 6,864,629 B2 6,867,539 B1 6,872,114 B2 6,872,248 B2 6,872,428 B2 6,897,474 B2 6,897,474 B2 6,997,607 B2 6,936,131 B2 6,975,067 B2 6,994,933 B1	11/2004 11/2004 12/2004 12/2004 1/2005 3/2005 3/2005 3/2005 3/2005 3/2005 5/2005	Affinito et al. Lee et al. Funkenbusch et al. Brown et al. Chung et al. Berard Miyaguchi et al. Burrows et al. McCormick et al. Chung et al. Mizutani et al. Yang et al. Chung et al. Weaver Silvernail et al. Duggal et al. Brown et al. Sugimoto et al. Komada Graff et al. McCormick et al. McCormick et al. Bates

7,015,640 B2	3/2006	Schaepkens et al.
7,018,713 B2	3/2006	Padiyath et al.
7,029,765 B2	4/2006	Kwong et al.
7,033,850 B2	4/2006	Tyan et al.
7,056,584 B2	6/2006	Iacovangelo
7,086,918 B2	8/2006	Hsiao et al.
7,122,418 B2	10/2006	Su et al.
7,156,942 B2	1/2007	McCormick et al.
7,166,007 B2	1/2007	Auch et al.
7,183,197 B2	2/2007	Won et al.
7,186,465 B2	3/2007	Bright
7,221,093 B2	5/2007	Auch et al.
7,255,823 B1	8/2007	Guenther et al.
2001/0015074 A1	8/2001	Hosokawa
2001/0015620 A1	8/2001	Affinito
2002/0022156 A1	2/2002	Bright
2002/0025444 A1	2/2002	Hebgrink et al.
2002/0068143 A1	6/2002	Silvernail et al.
2002/0069826 A1	6/2002	Hunt et al.
2002/0102363 A1	8/2002	Affinitio et al.
2002/0102818 A1	8/2002	Sandhu et al.
2002/0125822 A1	9/2002	Graff et al.
2002/0125022 A1	10/2002	Yamazaki et al.
2002/0139303 A1 2002/0140347 A1	10/2002	Weaver
2002/014034/ A1 2003/0038590 A1	2/2002	Silvernail et al.
	3/2003	
2003/0045021 A1		Akai
2003/0085652 A1	5/2003	Weaver
2003/0098647 A1	5/2003	Silvernail
2003/0117068 A1	6/2003	Forrest et al.
2003/0124392 A1	7/2003	Bright
2003/0127973 A1	7/2003	Weaver et al.
2003/0134487 A1	7/2003	Breen et al.
2003/0184222 A1	10/2003	Nilsson et al.
2003/0197197 A1	10/2003	Brown et al.
2003/0218422 A1	11/2003	Park et al.
2003/0235648 A1	12/2003	Affinito et al.
2004/0029334 A1	2/2004	Bijker et al.
2004/0046497 A1	3/2004	Schaepkens et al.
2004/0071971 A1	4/2004	Lacovangelo
2004/0113542 A1	6/2004	Hslao et al.
2004/0115402 A1	6/2004	Schaepkens
2004/0115859 A1	6/2004	Murayama et al.
2004/0119028 A1	6/2004	McCormick et al.
2004/0175512 A1	9/2004	Schaepkens
2004/0175580 A1	9/2004	Schaepkens
2004/0209090 A1	10/2004	Iwanaga
2004/0219380 A1	11/2004	Naruse et al.
2004/0229051 A1	11/2004	Schaepkens et al.
2004/0221454 A1	12/2004	Shaw et al.
2004/0263038 A1	12/2004	Ribotzi et al.
2005/0003098 A1		Kohler et al.
2005/0005098 A1 2005/0006786 A1	1/2005	Sawada
2005/0051094 A1		Schaepkens et al.
2005/0079295 A1	4/2005	Schaepkens
2005/0079380 A1		Iwanaga
2005/0093001 A1		Liu et al.
2005/0093437 A1		Ouyang
2005/0094394 A1	5/2005	Padiyath et al.
2005/0095422 A1		Sager et al.
2005/0095736 A1		Padiyath et al.
2005/0112378 A1		Naruse et al.
2005/0115603 A1	6/2005	Yoshida et al.
2005/0122039 A1		Satani
2005/0129841 A1	6/2005	McCormick et al.
2005/0133781 A1	6/2005	Yan et al.
2005/0140291 A1	6/2005	Hirakata et al.
2005/0146267 A1		Lee et al.
2005/0174045 A1		Lee et al.
2005/0202646 A1		Burrows et al.
2005/0212419 A1	9/2005	Vazan et al.
2005/0224935 A1		Schaepkens et al.

2005/02	238846 A1 10/2005	Arakatsu et al.	JP	09-161967		6/1997		
		Kim et al.	JP	9-161967		6/1997		
	003474 A1 1/2006		JP	9-201897		8/1997		
		Ohkubo	JP	09-232553		9/1997		
		McCormick et al.	JP	10-725		1/1998		
		Padiyath et al.	JР	10-013083		1/1998		
		McCormick et al.	JP	10-015085		1/1998		
		Furukawa et al.	JP	H10-41067		2/1998		
		Winters et al.	JP	10-334744		12/1998		
		Cok et al.	Л	11-017106		1/1999		
		Patry et al.	JP	11-040344		2/1999		
			Л					
		Okubo et al.		11-149826		6/1999		
2007/02	281089 A1 12/2007	Heller et al.	JP	11-255923		9/1999		
	FODEICN DATE		ЛР	2000-058258		2/2000		
	FOREIGN PATE	NT DOCUMENTS	JP	2002/505969		2/2002		
EP	0 390 540	10/1990	ЛР	10312883		3/2002		
EP	0 468 440 A2	1/1992	JP	2003/282239		10/2003		
EP	0 408 440 A2 0 547 550	6/1993	Л	3579556	B2	10/2004		
EP	0 547 550 A1	6/1993	JP	2006-294780		10/2006		
			WO	WO 87/07848		12/1987		
EP	0 590 467	4/1994	WO	WO 89/00337		1/1989		
EP	0 590 467 A1	4/1994	WO	WO 95/10117		5/1995		
EP	0 722 787	7/1996	WO	WO 96/23217		8/1996		
EP	0 722 787 A2	7/1996	WO	WO 97/04885		2/1997		
EP	0 777 280 B1	6/1997	WO	WO 97/16053		5/1997		
EP	0 777 280 A2	6/1997	WO	WO 97/22631		6/1997		
EP	0 777 281 A2	6/1997	WO	WO 98/10116		3/1998		
EP	0 787 824 A2	8/1997	WO	WO 98/18852		5/1998		
EP	0 787 826 A1	8/1997	WO	WO 99/16557		4/1999		
\mathbf{EP}	0 787 826	8/1997	WO	WO 99/16931		4/1999		
EP	0 915 105 A1	5/1999	WO	WO 99/46120		9/1999		
EP	0 916 394 A2	5/1999	WO	WO 00/26973		5/2000		
EP	0 916 394	5/1999	WO	WO 00/35603		6/2000		
EP	0 931 850 A1	7/1999	WO	WO 00/35604		6/2000		
EP	0 931 850	7/1999	WO	WO 00/35993		6/2000		
EP	0 977 469	2/2000	WO	WO 00/36661		6/2000		
EP	0 977 469 A2	2/2000	wo	WO 00/36665		6/2000		
EP	1 021 070 A1	7/2000	wo	00/53423	A 1	9/2000		
EP	1127 381 A1	8/2001	WO	WO 01 68360		9/2001		
EP	1 130 420 A2	9/2001	wo	WO 01/68360		9/2001		
EP	1 278 244 A2	1/2003	WO	WO 01/81649		11/2001		
EP	1 426 813 A2	6/2004	WO	WO 01/81049 WO 01/82336		11/2001		
EP	1 514 317 A1	3/2005	WO	WO 01/82389		11/2001		
GB	2 210 826 A	6/1989	WO	WO 01/82389 WO 01/87825		11/2001		
JP	S63-96895	4/1988	WO	WO 01/8/825 WO 01/89006		11/2001		
ЛЬ	63136316	8/1988	WO	WO 01/89000 WO 02/26973		4/2002		
JP	64-18441	1/1989	WO	WO 02/20973 WO 03/016589		2/2002		
JP	S64-41192	2/1989	WO					
JP	2-183230	7/1990		WO 03/098716		11/2003		
JP	3-183759	8/1991	WO WO	WO 2004/006199 WO 2004/016992		1/2004 2/2004		
JP	3-290375	12/1991						
JP	4-14440	1/1992	WO WO	WO 2004/070840		8/2004		
JP	4-48515	2/1992		WO 2004/089620		10/2004		
JP	4-1440	4/1992	WO	2004/112165		12/2004		
JP	H4-267097	9/1992	WO	WO 2005/015655		2/2005		
JP	05-217158	1/1993	WO	WO 2005/045947		5/2005		
JP	5-147678	6/1993	WO	WO 2005/048368		5/2005		
JP	H5-182759	7/1993	WO	2005/050754		6/2005		
JP	61-36159	5/1994	WO	WO 2006/036492	AI	4/2006		
JP	06158305	6/1994	WO	2008/140313		11/2008		
JP	61-79644	6/1994	WO	2008/142645		11/2008		
JP		8/1994		OTHER	PUE	BLICATION	IS	
JP JP	6-234186	3/1995	No					
JP JP	07-074378	6/1995		berg, H. et al.; Con				
	H07-147189			Through Polymers :				
JP ID	H7-192868	7/1995		, 2000; pp. 347–35				
JP ID	8-72188	3/1996	Yializ	is, A. et al.; Ultra	Higl	h Barrier Fi	lms; Denver.	, Apr.
JP ID	H8-171988	7/1996		, 2000; pp. 404–40				
JP	8-179292	7/1996		n, J.K. et al.; Requ				
JP	08325713	10/1996		ic Light Emitting I				
JP	8-318590	12/1996	ontine	s; Society of Vacu		Contorna Annd	Annual Ta-1	appn-
JP D	09059763	4/1997						nnear
JP	H9-132774	5/1997	Confe	rence Proceedings;	; Apr.	1999; pp. 4	130-439.	

Henry, B.M. et al.; Microstructural and Gas Barrier Properties of Transparent Aluminum Oxide and Indium Tim Oxide Films; Denver, Apr. 15–20, 2000; pp. 373–378; Society of Vacuum Coaters.

Affinito, J.D. et al.; Vacuum Deposited Polymer/metal Multilayer Films for Optical Applications; Paper No. C1.13; International Conference on Metallurgical Coatings; Apr. 15–21, 1995; pp. 1–14.

Affinito, J.D. et al.; Vacuum Deposition of Polymer Electrolytes On Flexible Substrates; The Ninth International Conference on Vacuum Web Coating; 1995; pp. 20–37.

Affinito, J.D. et al.; Vacuum Deposition of Polymer Electrolytes On Flexible Substrates; The Ninth International Conference on Vacuum Web Coating; 1995; pp. 0–16.

Affinito, J.D. et al.; Molecularly Doped Polymer Composite Films for Light Emitting Polymer Application Fabricated by the PML Process; 41st Technical Conference of the Society of Vacuum Coaters; Apr. 1998; pp. 220–225.

Affinito, J.D. et al., PML/Oxide/PML Barrier Layer Performance Differences Arising From Use OF UV Or Electron Beam Polymerization Of The PML Layers, SVC 40th Annual Technical Conference, Apr. 12–17, 1997, pp. 19–25.

Affinito, J.D. et al.; Polymer/Polymer, Polymer/Oxide, and Polymer/Metal Vacuum Deposited Interference Filters; Tenth International Vacuum Web Coating Conference; Nov. 1996; pp. 0–14.

Felts, J.T.; Transparent Barrier Coatings Update: Flexible Substrates; Society of Vacuum Coaters; 36th Annual Technical Conference Proceedings; Apr. 25–30, 1993; pp. 324–331.

Affinito, J.D. et al.; Ultra High Rate, Wide Area, Plasma Polymerized Films from High Molecular Weight/Low Vapor Pressure Liquid or Solid Monomer Precursors; 45th International Symposium of the American Vacuum Society; Nov. 2–6, 1998; pp. 0–26.

Tropsha et al.; Combinatorial Barrier Effect of the Multilayer SiOx Coatings on Polymer Substrates; 1997 Society of Vacuum Coaters; 40th Annual Technical Conference Proceedings; Apr. 12–17, 1997; pp. 64–69.

Tropsha et al., Activated Rate Theory Treatment of Oxygen and Water Transport through Silicon Oxide/Poly(ethylene terphthalate) Composite Barrier Structures; J. Phys. Chem B Mar. 1997; pp. 2259–2266.

Affinito, J.D. et al.; Vacuum Deposited Conductive Polymer Films; The Eleventh International Conference on Vacuum Web Coating; Nov. 9–11, 1997; pp. 0–12.

De Gryse, R. et al.; Sputtered Transparent Barrier Layers, Tenth International Conference on Vacuum Web Coating, Nov. 1996, pp. 190–198.

Hibino, N. et al.; Transparent Barrier $A1_20_3$ Coating By Activated Reactive Evaporation; Thirteenth International Conference on Vacuum Web Coating; Oct. 17–19, 1999; pp. 234–245.

Kukla, R. et al.; Transparent Barrier Coatings with EB–Evaporation, an Update; Section Five; Transparent Barrier Coating Papers; Thirteenth International Conference on Vacuum Web Coating; Oct. 17–19, 1999; pp. 222–233.

Bright, Clark I.; Transparent Barrier Coatings Based on ITO for Flexible Plastic Displays; Thirteenth International Conference on Vacuum Web Coating; Oct. 17–19, 1999; pp. 247–255.

Henry, B.M. et al.; Microstructural Studies of Transparent Gas Barrier Coatings on Polymer Substrates; Thirteenth International Conference on Vacuum Web Coating; Oct. 17–19, 1999; pp. 265–273.

Affinito, J.D. et al.; Ultra High Rate, Wide Area, Plasma Polymerized Films from High Molecular Weight/Low Vapor Pressure Liquid or Liquid/Solid Suspension Monomer Precursors; MRS Conference; Nov. 29–Dec. 3, 1998; Paper No. Y12.1

Affinito, J.D. et al., "Molecularly Doped Polymer Composite Films for Light Emitting Polymer Applications Fabricated by the PML Process" 41st Technical Conference of Society of Vacuum Coaters, Apr. 1998, pp. 1–6.

Affinito, J.D. et al., "Vacuum Deposition of Polymer Electrolytes on Flexible Substrates" The Ninth International Conference on Vacuum Web Coating, pp. 0–16.

Bunshah, R.F. et al., "Deposition Technologies for Films and Coatings" Noyes Publications, Park Ridge, New Jersey, 1982, p. 339.

Affinito, J.D., Energy Res. Abstr. 18(6), #17171, 1993.

Graupner, W. et al.; "High Resolution Color Organic Light Emitting Diode Microdisplay Fabrication Method", SPIE Proceedings, Nov. 6, 2000; pp. 11–19.

Czeremuszkin, G. et al.; Permeation Through Defects in Transparent Barrier Coated Plastic Films; 43rd Annual Technical Conference Proceedings; Apr. 15, 2000; pp. 408–413.

Affinito, J.D. et al.; Vacuum Deposited Conductive Polymer Films; The Eleventh International Conference on Vacuum Web Coatings, pp. 1–12.

Vossen, J.L. et al.; Thin Film Porcesses; Academic Press, 1978, Part II, Chapter II–1, Glow Dischareg Sputter Deposition, pp. 12–63; Part IV, Chapter IV–1 Plasma Deposition of Inorganic Compounds and Chapter IV–2 Glow Discharge Polymerization, pp. 335–397.

Affinito, J.D. et al.; Ultra High Rate, Wide Area, Plasma Polymerized Films from High Molecular Weight/Low Vapor Pressure Liquid or Solid Monomer Precursors; 45th International Symposium of the American Vacuum Society; pp. 0–26.

G. Gustafason, et al.; Flexible light–emitting diodes made from soluble conducting polymers; Letters to Nature; vol. 357; Jun. 11, 1992; pp. 477–479.

Tropsha et al.; Combinatorial Barrier Effect of the Multilayer SiOx Coatings on Polymer Substrates; 1997 Society of Vacuum Coaters, 40th Annual Technical Conferences Proceedings; pp. 64–69.

Tropsha et al.; Activated Rate Theory Treatment of Oxygen and Water Transport through Silicon Oxide/Poly(ethylene terphthalate) Composite Barrier Structures; J. Phys. Chem B 1997 pp. 2259–2266.

F.M. Penning; Electrical Discharges in Gases; 1965; pp. 1–51; Gordon and Breach, Science Publishers, New York–London–Paris.

Affinito, J.D. et al.; High Rate Vacuum Deposition of Polymer Electrolytes; Journal Vacuum Science Technology A 14(3), May/Jun. 1996.

Affinito, J.D. et al.; Vacuum Deposited Polymer/metal Multilayer Films for Optical Applications; Paper No. C1.13; pp. 1–14.

Shi, M.K. et al.; Plasma treatment of PET and acrylic coating surfaces–I. In situ XPS measurements; Journal of Adhesion Science and Technology; Mar. 2000 14(12); pp. 1–8.

Affinito, J.D. et al.; Vacuum Deposition of Polymer Electrolytes On Flexible Substrates, The Ninth International Conference on Vacuum Web Coating; pp. 20–37.

Affinito, J.D. et al.; Ultrahigh Rate, Wide Area, Plasma Polymerized Films from High Molecular Weight/Low Vapor Pressure Liquid or Solid Monomer Precursors; Journal Vacuum Science Technology A 17(4); Jul./Aug. 1999; pp. 1974–1981; American Vacuum Society.

Shi, M.K. et al.; In situ and real-time monitoring of plasma-induced etching PET and acrylic films, Plasmas and Polymers; Dec. 1999, 494); pp. 1–25.

Affinito, J.D. et al.; Vacuum Deposited Conductive Polymer Films; The Eleventh International Conference on Vacuum Web Coating; pp. 0–12.

Affinito, J.D. et al.; Molecularly Doped Polymer Composit Films for Light Emitting Polymer Application Fabricated by the PML Process; 41st Technical Conference of the Society of Vacuum Coaters; 1998; pp. 220–225.

Afffinto, J.D. et al.;Polymer/polymer, Polymer/Oxide, and Polymer/Metal Vacuum Deposited Interference Filters; Tenth International Vacuum Web Coating Conference; pp. 0–14.

Affinto, J.D. et al.; Vacuum Deposited Polymer/Metal Multilayer Films for Optical Application; Thin Solid Films 270, 1995; pp. 43–48.

Felts, J.T.; Transparent Barrier Coatings Update: Flexible Substrates; pp. 324–331.

Mahon, J.K., et al.; Requirements of Flexible Substrates for Organic Light Emitting Devices in Flat Panel Display Applications, Society of Vacuum Coaters, 42nd Annual Technical Conference Proceedings, 1999, pp. 456–459.

Henry, B.M. et al.; Microstructural and Gas Barrier Properties of Transparent Aluminium Oxide and Indium Tin Oxide Films; 2000; pp. 373–378; Society of Vacuum Coaters.

Phillips, R.W.; Evaporated Dielectric Colorless Films on PET and Opp Exhibiting High Barriers Toward Moisture and Oxygen; Society of Vacuum Coaters; 36th Annual Technical Conference Proceedings; 1993; pp. 293–300.

Yamada, Y. et al.; The Properties of a New Transparent and Colorless Barrier Film; 1995; pp. 28–31; Society of Vacuum Coaters.

Chahroudi, D.; Transparent Glass Barrier Coatings for Flexible Film Packaging; 1991; pp. 130–133; Society of Vacuum Coaters. Bright, Clark, I.; Transparent Barrier Coatings Based on ITo for Flexible Plastic Displays; pp. 247–255.

Henry, B.M. et al.; Microstructural Studies of Transparent Gas Barrier Coatings on Polymer Substates; pp. 265–273.

Hibino, N. et al.; Transparent Barrier Al/203 Coating By Activated Reactive Evaporation; pp. 234–245.

Kukla, R. et al.; Transparent Barrier Coatings with EB–Evaporation, an Update; Section Five; Transparent Barrier Coating Papers; pp. 222–233.

Krug, T. et al.; New Developments in Transparent Barrier Coatings; 1993; pp. 302–305; Society Vacuum Coaters.

Affinto, J.D. et al.; PML/Oxide/PML Barrier Layer Performance Differences Arising From Use Of UV or Electron Beam Polymerization of the PML Layers; Thin Solid Films; Elsevier Science S.A.; vol. 308–309; Oct. 31, 1997; pp. 19–25.

Affinito, J.D. et al.; A new method for fabricating transparent barrier layers, Thin Solid Films 290–291; 1996; pp. 63–67.

Affinito, J.D. et al.; Polymer–Oxide Transparent Barrier Layers; SVC 39th Annual Technical Conference; Vacuum Web Coating Session; 1996; pp. 392–397.

Hoffmann, G. et al.; Transparent Barrier Coatings by Reactive Evaporation; 1994; pp. 155–160; Society of Vacuum Coaters.

Norenberg, H. et al.; Comparative Study of Oxygen Permeation Through Polymers and Gas Barrier Films; 2000; pp. 347–351; Society of Vacuum Coaters.

Yializis, A. et al.; Ultra High Barrier Films; 2000; pp. 404–407; Society Vacuum Coaters.

Klemberg–Sapieha, J.E. et al.; Transparent Gas Barrier Coatings Produced by Dual–Frequency PECVD; 1993; pp. 445–449; Society of Vacuum Coaters.

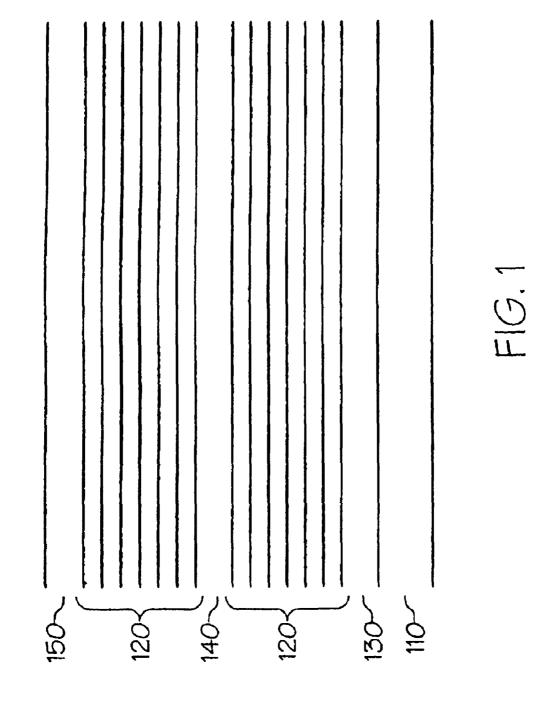
Finson, E. et al.; Transparent SiO2 Barrier Coatings: Conversion and Production Status; 1994; pp. 139–143; Society of Vacuum Coaters.

Yializis, A. et al.; High Oxygen Barrier Polypropylene Films Using Transparent Acrylate–A2O3 and Opaque Al–Acrylate Coatings; 1995; pp. 95–102; Society of Vacuum Coaters.

Shaw, D.G. et al.; Use of Vapor Deposited Acrylate Coatings to Improve the Barrier Properties of MetalLized Film; 1994; pp. 240–244; Society of Vacuum Coaters.

Wong, F.L., et al., "Long–lifetime thin–film encapsulated organic light–emitting diodes," Journal of Applied Physics 104, pp. 014509–1–4 (2008).

* cited by examiner



MULTILAYER PLASTIC SUBSTRATES

Matter enclosed in heavy brackets [] appears in the original patent but forms no part of this reissue specification; matter printed in italics indicates the additions 5 made by reissue.

This application is a continuation-in-part of U.S. patent application Ser. No. 09/427,138, filed Oct. 25, 1999, entitled "Environmental Barrier Material For Organic Light Emit- 10 ting Device and Method Of Making," now U.S. Pat. No. 6,522,067, issued Feb. 18, 2003.

BACKGROUND OF THE INVENTION

The present invention relates generally to plastic substrates which may be useful in products including, but not limited to, visual display devices, and more particularly to multilayer plastic substrates having improved light transmittance.

As used herein, the term "(meth)acrylic" is defined as "acrylic or methacrylic." Also, (meth)acrylate is defined as "acrylate or methacrylate."

As used herein, the term "average visible light transmittance" means the average light transmittance over the visible 25 range from 400 to 800 nm.

As used herein, the term "peak visible light transmittance" means the peak light transmittance over the visible range from 400 to 800 nm.

As used herein, the term "polymer precursor" includes ³⁰ monomers, oligomers, and resins, and combinations thereof. As used herein, the term "monomer" is defined as a molecule of simple structure and low molecular weight that is capable of combining with a number of like or unlike molecules to form a polymer. Examples include, but are not limited to, simple acrylate molecules, for example, hexanedioldiacrylate, or tetraethyleneglycoldiacrylate, styrene, methyl styrene, and combinations thereof. The molecular weight of monomers is generally less than 1000, while for fluorinated monomers, it is generally less than 2000. Monomers may be combined to form oligomers and resins but do not combine to form other monomers.

As used herein, the term "oligomer" is defined as a compound molecule of at least two monomers that maybe cured by radiation, such as ultraviolet, electron beam, or x-ray, glow discharge ionization, and spontaneous thermally induced curing. Oligomers include low molecular weight resins. Low molecular weight is defined herein as about 1000 to about 20,000 exclusive of fluorinated monomers. Oligomers are usually liquid or easily liquifiable. Oligomers do not combine to form monomers.

As used herein, the term "resin" is defined as a compound having a higher molecular weight (generally greater than 20,000) which is generally solid with no definite melting 55 point. Examples include, but are not limited to, polystyrene resins, epoxy polyamine resins, phenolic resins, and acrylic resins (for example, polymethylmethacrylate), and combinations thereof.

There is a need for versatile visual display devices for 60 electronic products of many different types. Although many current displays use glass substrates, manufacturers have attempted to produce commercial products, primarily liquid crystal display devices, using unbreakable plastic substrates. These attempts have not been completely successful to date 65 because of the quality, temperature, and permeation limitations of polymeric materials. Flexible plastic substrates,

such as polyethylene terephthalate (PET), polyethylene naphthalate (PEN), and polyether sulfone (PES), have been used in thicknesses from about 0.004 inches to 0.007 inches. However, the surface quality of these substrates is often poor, with the surface having large numbers of scratches, digs, pits, and other defects.

In addition, many polymers exhibit poor oxygen and water vapor permeation resistance, often several orders of magnitude below what is required for product performance. For example, the oxygen transmission rates for materials such polyethylene terephthalate (PET) are as high as 1550 $\text{cc/m}^2/\text{day/micron}$ of thickness (or 8.7 $\text{cc/m}^2/\text{day}$ for 7 mil thickness PET), and the water vapor transmission rates are also in this range. Certain display applications, such as those using organic light emitting devices (OLEDs), require encapsulation that has a maximum oxygen transmission rate of 10^{-4} to 10^{-2} cc/m²/day, and a maximum water vapor transmission rate of 10^{-5} to 10^{-6} g/m²/day.

Barrier coatings have been applied to plastic substrates to decrease their gas and liquid permeability. Barrier coatings typically consist of single layer thin film inorganic materials, such as Al, SiO_x , AlO_x , and Si_3N_4 vacuum deposited on polymeric substrates. A single layer coating on PET reduces oxygen permeability to levels of about 0.1 to 1.0 cc/m²/day, and water vapor permeability of about 0.1 to 1.0 g/m²/day. However, those levels are still insufficient for many display devices.

Additionally, many processes used in the manufacture of displays require relatively high temperatures that most polymer substrates cannot tolerate. For example, the recrystallization of amorphous Si to poly-Si in thin film transistors requires substrate temperatures of at least 160°-250° C., even with pulsed excimer laser anneals. The conductivity of a transparent electrode, which is typically made of indium tin oxide (ITO), is greatly improved if deposition occurs above 220° C. Polyimide curing generally requires temperatures of 250° C. In addition, many of the photolithographic process steps for patterning electrodes are operated in excess of 120° C. to enhance processing speeds in the fabrication. These processes are used extensively in the manufacture of display devices, and they have been optimized on glass and silicon substrates. The high temperatures needed for such processes can deform and damage a plastic substrate, and subsequently destroy the display. If displays are to be manufactured on flexible plastic materials, the plastic must be able to withstand the necessary processing conditions, including high temperatures over 100° C., harsh chemicals, and mechanical damage.

Thus, there is a need for an improved plastic substrate for visual display devices, and for a method of making such a substrate.

SUMMARY OF THE INVENTION

The present invention meets this need by providing a multilayer plastic substrate. The substrate consists essentially of a plurality of thin film layers of at least one polymer, the plurality of thin films layers being adjacent to one another and having sufficient strength to be self-supporting, wherein the multilayer plastic substrate has an average visible light transmittance of greater than about 80%. The average visible light transmittance is typically greater than about 85%, and it can be greater than about 90%. The peak visible transmittance is typically greater than about 85% and it can be greater than about 90%.

There are typically at least about 50 thin film layers. The number of layers depends on the thickness of the thin film

55

60

layers and the desired overall thickness of the multilayer plastic substrate. The multilayer plastic substrate is typically at least about 0.001 inches thick, and generally at least about 0.004 inches thick. Each thin film layer is typically less than about 50 μ m thick.

Polymers include, but are not limited to (meth)acrylatecontaining polymers, styrene containing polymers, methyl styrene containing polymers, and fluorinated polymers, and combinations thereof. The glass transition temperature of the at least one polymer is generally greater than about 150°¹⁰ C., and it may be greater than about 200° C.

The surface roughness of the multilayer plastic substrate is generally less than about 10 nm, and it may be less than about 5 nm, or less than about 2 nm.

The multilayer plastic substrate can have a refractive index of greater than about 1.4 or greater than about 1.5.

The multilayer plastic substrate can include additional layers, including, but not limited to, scratch resistant layers, antireflective coatings, antifingerprint coatings, antistatic 20 coatings, conductive coatings, transparent conductive coatings, and barrier coatings, to provide functionality to the substrate if desired.

Another aspect of the invention involves a method of making the multilayer plastic substrate. The method includes 25 providing a support, depositing a plurality of thin film layers of at least one polymer on the support so that the plurality of thin film layers have sufficient strength to be self-supporting to form the multilayer substrate, and removing the support from the multilayer substrate, wherein the multilayer plastic 30 substrate has an average visible light transmittance of greater than about 80%.

The thin film layers can be deposited in a vacuum. One example of a vacuum deposition process is flash evaporation. In this method, depositing the plurality of thin film ³⁵ layers includes flash evaporating a polymer precursor, condensing the polymer precursor as a liquid film, and cross-linking the polymer precursor to form the polymer. The polymer precursor can be cross-linked by any suitable method, including, but not limited to, radiation curing, such ⁴⁰ as ultraviolet, electron beam, or x-ray, glow discharge ionization, and spontaneous thermally induced curing.

Alternatively, the plurality of thin film layers can be deposited by extruding or casting a layer of polymer precursor, and cross-linking the polymer precursor to form ⁴⁵ the polymer using any suitable cross-linking method.

Accordingly, it is an object of the present invention to provide an improved, multilayer plastic substrate and to provide a method of making such a substrate.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. **1** is a cross-section of one embodiment of the substrate of the present invention.

DESCRIPTION OF THE INVENTION

FIG. 1 shows one embodiment of a multilayer plastic substrate of the present invention. The multilayer plastic substrate 100 is formed on a support 110. After the multilayer plastic substrate is formed, the support 110 is removed.

The multilayer plastic substrate of the present invention consists essentially of a plurality of thin film layers **120** of at least one polymer adjacent to one another. By adjacent, we mean next to, but not necessarily directly next to. In most of the multilayer plastic substrate, the polymer thin film layers ⁶⁵ will be directly next to one another. However, there can be additional layers intervening between some adjacent layers

in order to provide additional functionality to the multilayer plastic substrate, as shown in FIG. 1 and described below.

The plurality of thin film layers have sufficient strength to be self-supporting after they are formed. The exact number of thin film layers is not critical. It depends on the thickness of each of the individual thin film layers and the desired overall thickness of the multilayer plastic substrate. There must be enough thin film layers so that the plurality of thin film layers have sufficient strength to be self-supporting. As used herein, the term self-supporting means the substrate can be handled and processed without the need for an underlying support once the plurality of thin film layers have been deposited. There are typically at least about 50 thin film layers, more typically at least about 100 thin film layers. There are generally in the range of about 500 thin film layers to about 1000 thin film layers or more. Each thin film layer is typically between about 0.05 to about 2 µm thick, generally between about 0.2 to about 0.3 µm. If the thin film layers are extruded, they are usually thicker, typically up to about 50 um thick, in that case. The multilayer plastic substrate is typically at least about 0.001 inches thick, and generally at least about 0.004 inches thick. A 0.007 inch thick substrate would require about 90 to 350 passes of the web past the polymer precursor sources. The multilayer plastic substrate can be flexible or rigid.

The average visible light transmittance of the multilayer plastic substrate is greater than about 80%, generally greater than 85%, and it may be greater than 90%. The peak visible light transmittance is generally greater than 85%, and it may be greater than 90%.

The at least one polymer can be any suitable polymer, including, but not limited to, polymers made from styrene polymer precursors, polymers made from methyl styrene polymer precursors, for example, polymers made from hexanedioldiacrylate or tetraethyleneglycoldiacrylate polymer precursors, and fluorinated polymers, and combinations thereof. Polymers made from (meth)acrylate polymer precursors work well.

The multilayer plastic substrate can be flexible or rigid. Multilayer plastic substrates made from polymers including, but not limited to, (meth)acrylate polymer precursors will be flexible. One advantage of multilayer laminated materials is that they typically have greater strength and flexibility than comparable single layer substrates. A multilayer plastic substrate of the present invention generally has hundreds of cross-linked layers that provide mechanical strength and sufficient rigidity to support the circuitry and devices on the display.

A multilayer plastic substrate made from (meth)acrylate polymer precursors will have excellent transmission at visible wavelengths. Because polymers made from (meth) acrylate polymer precursors have very low optical absorption, a multilayer plastic substrate made entirely from such polymers will have high optical transparency, typically an average visible light transmittance of greater than about 90%. Multilayer substrates made entirely from fluorinated polymers will also have an average visible light transmittance of greater than 90%. Substrates made from styrene and methyl styrene polymers would have an average visible light transmittance of about 89%.

The birefringence present in many flexible substrates can be reduced or eliminated with the present invention because the multilayer plastic substrate is not mechanically stressed during deposition.

Fully cured layers of polymers made from (meth)acrylate polymer precursors generally have a refractive index of

25

65

greater than about 1.5, while fully cured fluorinated polymers generally have a refractive index of greater than about 1.4. Styrene containing polymers would have a refractive index of about 1.6.

Many optical applications, such as mirrors and reflectors, 5 and display applications, such as organic light emitting devices, require substrates with a surface roughness of less than 2 nm. Surface roughness is the root mean square of peak-to-valley measurement over a specified distance, usually 1 nm. It can be measured using an atomic force microscope or back reflection distribution function. Many substrates do not have the necessary surface smoothness. For example, the surface roughness of PET is about 20-50 nm with 100 nm spikes. In contrast, flash evaporated polymer coatings have a very low surface roughness, generally less than about 10 nm, and it may be less than 5 nm, or less than about 2 nm. Surface roughness on the order of 1 nm has been demonstrated. The surface of the multilayer plastic substrate is specular because of the exceptional smoothness of the polymer layers. 20

Because the polymer material is highly cross-linked, the multilayer plastic substrate can have a high glass transition temperature and excellent chemical resistance. The glass transition temperature of the at least one polymer is generally greater than about 150° C., and may be greatr than about 200° C.

Polymers including, but not limited to, (meth)acrylates, polycarbonates, polysulfones, polyethersulfones, polymides, polyamides, and polyether napthteates have demonstrated excellent resistance to solvents. This provides protection from processing chemicals, ultraviolet light exposure, and photoresists during lithography processes used to manufacture flat panel displays and their devices.

The thin film layers that form the multilayer substrate can be deposited by any suitable method, including vacuum flash 35 evaporation, extrusion, or casting. With vacuum flash evaporation, deposition can be performed using a rotating drum or strap configuration. The polymer precursor is degassed and metered into a hot tube where it flash evaporates and exits through a nozzle as a polymer precursor gas. 40

The flash evaporating may be performed by supplying a continuous liquid flow of the polymer precursor into a vacuum environment at a temperature below both the decomposition temperature and the polymerization temperature of the polymer precursor, continuously atomizing the 45 polymer precursor into a continuous flow of droplets, and continuously vaporizing the droplets by continuously contacting the droplets on a heated surface having a temperature at or above a boiling point of the liquid polymer precursor, but below a pyrolysis temperature, forming the evaporate. $_{50}$ The droplets typically range in size from about 1 micrometer to about 50 micrometers, by they could be smaller or larger.

Alteratively, the flash evaporating may be performed by supplying a continuous liquid flow of the polymer precursor into a vacuum environment at a temperature below both the 55 decomposition temperature and the polymerization temperature of the polymer precursor, and continuously directly vaporizing the liquid flow of the polymer precursor by continuously contacting the liquid polymer precursor on a heated surface having a temperature at or above the boiling 60 point of the liquid polymer precursor, but below the pyrolysis temperature, forming the evaporate. This may be done using the vaporizer disclosed in U.S. Pat. Nos. 5,402,314, 5,536,323, and 5,711,816, which are incorporated herein by reference.

The polymer precursor then condenses on the support as a liquid film which is subsequently cross-linked to form a

6

polymer by any suitable method, including, but not limited to, radiation, such as ultraviolet, electron beam, or x-ray, glow discharge ionization, and spontaneous thermally induced curing. This process is capable of depositing thousands of polymer layers at web speeds up to 100 m/min.

Alteratively, after degassing, the polymer precursor can be deposited by extruding, spraying, or casting layers of polymer precursor on the support. The polymer precursor is then cross-linked using any suitable method, such as those described above.

The functionality of the multilayer plastic substrate can be increased by the incorporation of functional layers 130, 140, and 150 during the deposition process. These functional layers 130, 140, and 150 can be deposited at any time during the deposition process. They can be deposited below, 130, in between, 140, or on top of, 150, the plurality of thin film layers 120 of the multilayer plastic substrate, as shown in FIG. 1. As used herein, depositing a coating adjacent to the multilayer plastic substrate includes: depositing the coating on the top layer of the multilayer plastic coating; depositing the coating on the multilayer plastic substrate and then depositing additional layers of the multilayer plastic substrate over the coating so that the coating is between the layers of the multilayer plastic substrate; and depositing the coating first and then depositing the layers of the multilayer plastic substrate, and combinations thereof. Functional layers 130, 140, and 150 include, but are not limited to, scratch resistant coatings, antirefelctive coatings, antifingerprint coatings, antistatic coatings, conductive coatings, transparent conductive coatings, and barrier coatings, and other functional layers. Depositing these additional layers allows the multilayer plastic substrate to be specifically tailored to different applications. Little or no surface modification is necessary for deposition of other layers because of the very smooth surface of the multilayer plastic substrate. Interfaces can be graded to bond all integrated functional layers firmly during the same coating run and pumpdown.

For some applications, it may be important that the presence of functional layers not reduce the average visible light transmittance below 80%, for others, not below 85%, and still others, not below 90%. In others, it may be important that the peak visible light transmittance not drop below 85%, and for others, not below 90%. In others, it may be important that the functional layers not increase the surface roughness to greater than about 10 nm, for others, not greater than about 5 nm, and for others, not greater than 2 nm.

One type of functional layer that can be included is a barrier coating. One example of a barrier coating is described in application Ser. No. 09/427,138, filed Oct. 25, 1999, entitled "Environmental Barrier Material for Organic Light Emitting Device and Method of Making," which is incorporated herein by reference. The barrier coating can be a barrier stack having one or more barrier layers and one or more polymer layers. There could be one polymer layer and one barrier layer, there could be one or more polymer layers on one side of one or more barrier layers, or there could be one or more polymer layers on both sides of one or more barrier layers. The important feature is that the barrier stack have at least one polymer layer and at least one barrier layer. The barrier layers and polymer layers in the barrier stack can be made of the same material or of a different material. The barrier layers are typically in the range of about 100-400 Å thick, and the polymer layers are typically in the range of about 1000-10,000 Å thick.

The number of barrier stacks is not limited. The number of barrier stacks needed depends on the material used for the

polymer of the substrate and the level of permeation resistance needed for the particular application. One or two barrier stacks should provide sufficient barrier properties for some applications. The most stringent applications may require five or more barrier stacks.

The barrier layers should be transparent. Transparent barrier materials include, but are not limited to, metal oxides, metal nitrides, metal carbides, metal oxynitrides, metal oxyborides, and combinations thereof. The metal oxides include, but are not limited to, silicon oxide, aluminum 10 oxide, titanium oxide, indium oxide, tin oxide, indium tin oxide, tantalum oxide, zirconium oxide, niobium oxide, and combinations thereof. The metal carbides include, but are not limited to, boron carbide, tungsten carbide, silicon carbide, and combinations thereof. The metal nitrides include, but are not limited to, aluminum nitride, silicon nitride, boron nitride, and combinations thereof. The metal oxynitrides include, but are not limited to, aluminum oxynitride, silicon oxynitride, boron oxynitride, and combinations thereof. The metal oxyborides include, but are not 20 limited to, zirconium oxyboride, titanium oxyboride, and combinations thereof.

The polymer layers of the barrier stacks can be made from (meth)acrylate polymer precursors. The polymer layers in the barrier stacks can be the same or different.

The barrier stacks can be made by vacuum deposition. ²⁵ The barrier layer can be vacuum deposited onto, or into, the multilayer plastic substrate, or another functional layer. The polymer layer is then deposited on the barrier layer, preferably by flash evaporating (meth)acrylate polymer precursors, condensing on the barrier layer, and polymerizing in situ in a vacuum chamber. U.S. Pat. Nos. 5,440,446 and 5,725,909, which are incorporated herein by reference, describe methods of depositing thin film, barrier stacks.

Vacuum deposition includes flash evaporation of (meth) acrylate polymer precursors with in situ polymerization ³⁵ under vacuum, plasma deposition and polymerization of (meth)acrylate polymer precursors, as well as vacuum deposition of the barrier layers by sputtering, chemical vapor deposition, plasma enhanced chemical vapor deposition, evaporation, sublimation, electron cyclotron resonanceplasma enhanced vapor deposition (ECR-PECVD), and combinations thereof.

In order to protect the integrity of the barrier layer, the formation of defects and/or microcracks in the deposited layer subsequent to deposition and prior to downstream pro-45 cessing should be avoided. The multilayer plastic substrate is preferably manufactured so that the barrier layers are not directly contacted by any equipment, such as rollers in a web coating system, to avoid defects that may be caused by abrasion over a roll or roller. This can be accomplished by 50 designing the deposition system such that the barrier layers are always covered by polymer layers prior to contacting or touching any handling equipment.

While certain representative embodiments and details have been shown for purposes of illustrating the invention, it will be apparent to those skilled in the art that various changes in the compositions and methods disclosed herein may be made without departing from the scope of the invention, which is defined in the appended claims.

What is claimed is:

1. A multilayer plastic substrate consisting essentially of:

a plurality of flash evaporated thin film layers of at least one polymer, the plurality of thin film layers being adjacent to one another and having sufficient strength to be self-supporting, wherein the multilayer plastic substrate has an average visible light transmittance of greater than about 80%, wherein the multilayer plastic substrate comprises at least about 50 thin film layers, and wherein the multilayer plastic substrate has a surface roughness of less than about 10 nm.

2. The multilayer plastic substrate of claim **1** wherein the average visible light transmittance is greater than about 85%.

3. The multilayer plastic substrate of claim **1** wherein the average visible light transmittance is greater than about 90%.

4. The multi layer plastic substrate of claim **1** wherein the peak visible light transmittance is greater than about 85%.

5. The multilayer plastic substrate of claim 1 wherein the peak visible light transmittance is greater than about 90%.

6. The multilayer plastic substrate of claim **1**, wherein the multilayer plastic substrate comprises at least about 100 thin film layers.

7. The multilayer plastic substrate of claim 6, wherein the multilayer plastic substrate comprises at least about 500 thin film layers.

8. The multilayer plastic substrate of claim 7, wherein the multilayer plastic substrate comprises at least about 1000 thin film layers.

9. The multilayer plastic substrate of claim **1**, wherein the multilayer plastic substrate is at least about 0.001 inches thick.

10. The multilayer plastic substrate of claim **1**, wherein the multilayer plastic substrate is at least about 0.004 inches thick.

11. The multilayer plastic substrate of claim 1, wherein each thin film layer is less than about 50 μ m thick.

12. The multilayer plastic substrate of claim 1, wherein each thin film layer is less than about $5 \,\mu\text{m}$ thick.

13. The multilayer plastic substrate of claim 1, wherein each thin film layer is in the range of about 0.05 to about 2 μ m thick.

14. The multilayer plastic substrate of claim 1, wherein each thin film layer is in the range of about 0.2 to about 0.3 μ m.

15. The multilayer plastic substrate of claim **1**, wherein the at least one polymer is selected from (meth)acrylates, polystyrenes, methyl styrene-containing polymers, fluorinated polymers, polycarbonates, polysulfones, polyethersulfones, polyethersulfones, and polyether naphthalenes, and combinations thereof.

16. The multilayer plastic substrate of claim 1, wherein the glass transition temperature of the at least one polymer is greater than about 150° C.

17. The multilayer plastic substrate of claim 1, wherein the glass transition temperature of the at least one polymer is greater than about 200° C.

18. The multilayer plastic substrate of claim **1**, wherein the multilayer plastic substrate has a surface roughness of less than about 2 nm.

19. The multilayer plastic substrate of claim 1, wherein the multilayer plastic substrate has a refractive index of greater than about 1.5.

20. The multilayer plastic substrate of claim **1**, wherein the multilayer plastic substrate has a refractive index of greater than about 1.4.

21. The multilayer plastic substrate of claim **1**, wherein the multilayer plastic substrate is flexible.

22. The multilayer plastic substrate of claim 1, wherein the multilayer plastic substrate is rigid.

23. The multilayer plastic substrate of claim **1**, wherein the multilayer plastic substrate has a surface roughness of less than about 5 nm.

* * * * *