

### (12) United States Patent Zdeblick et al.

### (54) COMMUNICATION SYSTEM WITH PARTIAL POWER SOURCE

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#### (56)**References Cited**

### U.S. PATENT DOCUMENTS

1,548,459 A 8/1925 Hammer 2,587,158 A 2/1952 Hofberg (Continued)

### FOREIGN PATENT DOCUMENTS

1588649 3/2005 CN CN 1650844 8/2005 (Continued)

### OTHER PUBLICATIONS

AADE, "AADE 37th Annual Meeting San Antonio Aug. 4-7, 2010" American Association of Diabetes Educators (2010); http://www. diabeteseducator.org/annualmeeting/2010/index.html; 2 pp.

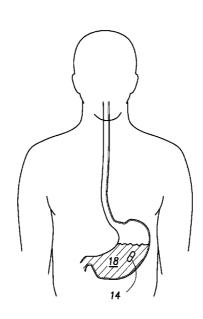
(Continued)

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### **ABSTRACT**

The system of the present invention includes a conductive element, an electronic component, and a partial power source in the form of dissimilar materials. Upon contact with a conducting fluid, a voltage potential is created and the power source is completed, which activates the system. The electronic component controls the conductance between the dissimilar materials to produce a unique current signature. The system can be used in a variety of different applications, including as components of ingestible identifiers, such as may be found in ingestible event markers, e.g., pharmainformatics enabled pharmaceutical compositions.

### 18 Claims, 3 Drawing Sheets



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See application file for complete search history.

### (56) References Cited

### U.S. PATENT DOCUMENTS

2,973,555	A	3/1961	Schwepke
3,048,526	A	8/1962	Boswell
3,079,824	A	3/1963	Schott
3,096,248	A	7/1963	Rudzki
3,176,399	A	4/1965	Marino et al.
3,589,943	A	6/1971	Grubb et al.
3,607,788	A	9/1971	Adolph
3,642,008	A	2/1972	Bolduc
3,679,480	A	7/1972	Brown et al.
3,682,160	A	8/1972	Murata
3,719,183	A	3/1973	Schwartz
3,799,802	A	3/1974	Schneble, Jr. et al.
3,828,766	A	8/1974	Krasnow
3,837,339	A	9/1974	Aisenberg et al.
3,849,041	A	11/1974	Knapp
3,893,111	A	7/1975	Cotter
3,944,064	A	3/1976	Bashaw et al.
3,967,202	A	6/1976	Batz
3,989,050	A	11/1976	Buchalter
4,017,856	A	4/1977	Wiegand
4,055,178	A	10/1977	Harrigan
4,062,750	A	12/1977	Butler
4,077,397	A	3/1978	Ellis
4,077,398	A	3/1978	Ellis
4,082,087	A	4/1978	Howson

5/1978 Long 8/1978 Auphan 4,090,752 A 4,106,348 A 12/1978 4,129,125 A Lester 2/1979 4,139,589 A Beringer et al. 4,143,770 A 3/1979 Grimmell et al. 4,166,453 A 9/1979 McClelland 4,239,046 A 12/1980 Ong 4,251,795 A 2/1981 Shibasaki et al. 4,269,189 A 5/1981 Abraham 4.331.654 A 5/1982 Morris 4.345,588 A 8/1982 Widder et al. 12/1983 4,418,697 Tama 1/1984 4,425,117 A Hugemann 4,439,196 A 3/1984 Higuchi 4,494,950 A 1/1985 Fischell 4,559,950 A 12/1985 Vaughan 4,564,363 A 1/1986 Bagnall et al. 4,635,641 A 1/1987 Hoffman 4,654,165 A 3/1987 Eisenber 4.663,250 A 5/1987 Ong et al. 4,669,479 A 6/1987 Dunseath 4,687,660 A 8/1987 Baker et al. 4.725.997 2/1988 Urquhart et al. 4,749,575 A 6/1988 Rotman et al. 4,763,659 A 8/1988 Dunseath 4,767,627 8/1988 Caldwell et al. 4,775,536 A 10/1988 Patell 4,784,162 A 11/1988 Ricks Benjamin et al. 4.793.825 A 12/1988 3/1989 4,814,181 A Jordan et al. 4,844,076 A 7/1989 Lesho 4,847,090 A 7/1989 Della Posta et al. 4,876,093 A 10/1989 Theeuwes et al. 4,891,223 A 1/1990 Ambegaonakar et al. 1/1990 4,896,261 A Nolan 4,900,552 A 2/1990 Sanvordeker et al. 4,975,230 A 12/1990 Pinkhasov 4,987,897 A 1/1991 Funke 3/1991 5.000.957 Eckenhoff et al. 5/1991 5,016,634 A Vock et al. 5,018,335 A 5/1991 Yamamoto et al. 5,079,006 A 1/1992 Urguhart Kinlen et al. 5,110,441 A 5/1992 5,160,885 A 11/1992 Hannam et al. 5,167,626 A 12/1992 Casper 5.176.626 A 1/1993 Soehendra 5,187,723 A 2/1993 Mueller 5/1993 5,213,738 A Hampton et al. 6/19935,218,343 A Stobbe et al 5,261,402 A 11/1993 DiSabito 5,263,481 A 11/1993 Axelgaard et al. 5,273,066 A 12/1993 Graham et al. 5,279,607 1/1994 Schentag et al. 5,281,287 A 1/1994 Lloyd 5.283,136 A 2/1994 Peled et al. 5,288,564 A 2/1994 Klein 5,305,745 A 4/1994 Zacouto 5/1994 5,310,301 A Aono 5,318,557 A 6/1994 Gross 5,331,953 A 7/1994 Andersson et al. 5,394,882 A 3/1995 Mawhinney 5,395,366 A 3/1995 D'Andrea et al. 5,436,091 A 7/1995 Shackle et al. 5,443,461 A 8/1995 Atkinson et al. 5,443,843 A 8/1995 Curatolo et al. 10/1995 5,458,141 A Neil et al. 5,458,994 A 10/1995 Nesselbeck et al. 5,485,841 A 1/1996 Watkin et al. 5,506,248 A 4/1996 Nikfar et al. 5,551,020 A 8/1996 Flax et al. 5,567,210 A 10/1996 Bates et al 1/1997 5.596,302 A Mastrocola et al. 5,600,548 A 2/1997 Nguyen et al. 5,603,363 A 2/1997 Nelson 5,634,468 A 6/1997 Platt

5,645,063 A

5,659,247 A

5,703,463 A

5,705,189 A

7/1997

8/1997

12/1997

Straka et al.

Clements

Smith

1/1998 Lehmann et al.

## US 10,542,909 B2

Page 3

(56)		Referen	ces Cited		6,602,518 6,605,038		8/2003 8/2003	Seielstad et al.
	U.S.	PATENT	DOCUMENTS		6,609,018	B2	8/2003	Cory
5 704 42	2 4	2/1009	Daywat at al		6,612,984 6,632,175		9/2003	Marshall
5,724,432 5,738,700			Bouvet et al. Peachey et al.		6,632,216	B2	10/2003	Houzego et al.
5,740,81	1 A	4/1998	Hedberg		6,635,279			Kolter et al.
5,757,320			Koyama et al.		6,643,541 6,654,638			Mok et al. Sweeney
5,772,573 5,792,043			Lesinski et al. Schaefer		6,663,846			McCombs
5,802,46			Salazar		6,673,474	B2		Yamamoto
5,833,710		11/1998			6,680,923 6,689,117		1/2004	Leon Sweeney et al.
5,842,324 5,845,265			Grosskopf et al. Woolston		6,694,161			Mehrotra
5,862,80		1/1999			6,704,602	B2		Berg et al.
5,868,130	5 A	2/1999			6,720,923			Hayward et al. Christophersom et al.
5,914,132	2 A *	6/1999	Kelm	A61K 9/145 424/467	6,738,671 6,740,033			Olejniczak et al.
5,914,70	1 A	6/1999	Gersheneld et al.	424/40/	6,745,082	B2	6/2004	Axelgaard et al.
5,925,030			Gross et al.		6,755,783			Cosentino
5,957,854			Besson et al.		6,757,523 6,759,968		6/2004 7/2004	
5,963,132 5,974,12			Yoakum et al. Schlueter, Jr. et al.		6,767,200	B2	7/2004	Sowden et al.
5,981,16			Mandecki		6,773,429			Sheppard et al.
5,999,840			Pardey et al.		6,800,060 6,801,137			Marshall Eggers et al.
6,018,229 6,038,464			Mitchell et al. Axelgaard et al.		6,816,794		11/2004	Alvi
6,042,710		3/2000	Dubrow		6,822,554			Vrijens et al.
6,047,203			Sackner		6,824,512 6,836,862			Warkentin et al. Erekson et al.
6,068,469 6,068,589			Wilson Neukermans		6,839,659		1/2005	Tarassenko et al.
6,076,010			Feierbach et al.		6,840,904			Goldberg
6,081,73		6/2000			6,842,636 6,845,272			Perrault Thomsen
6,091,97: 6,095,98:			Daddona et al. Raymond et al.		6,864,780		3/2005	
6,115,63		9/2000			6,879,810		4/2005	
6,122,35			Schlueter, Jr. et al.		6,888,337 6,889,165			Sawyers Lind et al.
6,141,592 6,149,940		10/2000	Pauly Maggi et al.		6,909,878		6/2005	Haller
6,200,26			Walsh et al.		6,922,592			Thompson et al.
6,206,70	2 B1		Hayden et al.		6,928,370 6,929,636			Anuzis et al. Von Alten
6,217,74 6,231,59			Crosby Meserol		6,937,150			Medema
6,245,05			Sieben et al.		6,942,616		9/2005	
6,269,05	8 B1	7/2001	Yamanoi et al.		6,942,770 6,951,536		9/2005 10/2005	Cai et al.
6,285,89° 6,287,252		9/2001 9/2001	Kilcoyne et al.		6,957,107			Rogers et al.
6,288,629			Cofino et al.		6,958,603		10/2005	Kondo
6,289,23	8 B1		Besson et al.		6,960,617 6,968,153			Omidian et al. Heinonen
6,315,719 6,317,714			Rode et al. Del Castillo					Patel et al.
6,342,77			Kreisinger et al.		6,982,094			Sowden
6,344,82	4 B1	2/2002	Takasugi et al.		6,987,965 6,990,082			Ng et al. Zehavi et al.
6,358,202 6,364,834		3/2002 4/2002			7,002,476			Rapchak
6,366,20			Ishikawa et al.		7,004,395			Koenck
6,371,92	7 B1	4/2002			7,009,634 7,009,946			Iddan et al. Kardach
6,374,670 6,380,853			Spelman Yarin et al.		7,003,340			Gorsuch
6,390,08			Noehl et al.		7,016,648		3/2006	
6,394,99	7 B1		Lemelson		7,020,508 7,024,248			Stivoric Penner et al.
6,426,863 6,432,293			Munshi Pinto et al.		7,024,246		4/2006	
6,440,069	2 B1		Raymond et al.		7,031,857	B2		Tarassenko et al.
6,441,74	7 B1	8/2002	Khair		7,039,453 7,044,911			Mullick Drinan et al.
6,453,199 6,477,424			Kobozev Thompson et al.		7,044,511			Awater et al.
6,496,70			Ng et al.		7,061,236	B2	6/2006	
6,526,31	5 B1	2/2003	Inagawa		7,083,578			Lewkowicz
6,531,020 6,544,17		3/2003 4/2003	Takeichi et al.		7,116,252 7,118,531		10/2006	Teraguchi Krill
6,547,99			Monkhouse et al.		7,122,143			Sowden et al.
6,564,079	9 B1	5/2003	Cory		7,127,300			Mazar et al.
6,567,683			Takamori et al.		7,146,228		12/2006	Nielsen Do et al.
6,572,630 6,577,893			Hagen et al. Besson		7,146,449 7,149,581			Goedeke
6,579,23		6/2003	Phipps		7,154,071		12/2006	Sattler et al.
6,595,929		7/2003	Stivoric		7,155,232	B2	12/2006	Godfrey et al.
6,599,284	4 B2	7/2003	Faour et al.		7,160,258	В2	1/2007	Imran

## US 10,542,909 B2

Page 4

(56)	Referen	ces Cited	7,525,426 7,537,590			Edelstein
U	.S. PATENT	DOCUMENTS	7,539,533	B2	5/2009 5/2009	Tran
= 464040 D			7,542,878 7,547,278		6/2009 6/2009	Nanikashvili Miyazaki et al.
7,164,942 B 7,171,166 B		Avrahami Ng et al.	7,551,590		6/2009	
7,171,177 B	2 1/2007	Park et al.	7,554,452		6/2009	
7,171,259 B 7,176,784 B		Rytky Gilbert et al.	7,558,620 7,575,005			Ishibashi Mumford
7,170,784 B			7,616,111	B2	11/2009	Covannon
7,188,199 B		Leung et al.	7,617,001 7,626,387		11/2009 12/2009	Penner et al.
7,188,767 B 7,194,038 B		Penuela Inkinen	7,639,473	B2	12/2009	Hsu et al.
7,196,495 B	3/2007	Burcham	7,640,802			King et al.
7,206,630 B 7,209,790 B		Tarler Thompson et al.	7,645,262 7,647,112	B2	1/2010	Greenberg et al. Tracey
7,215,660 B	2 5/2007	Perlman	7,647,185	B2	1/2010	Tarassenko et al.
7,215,991 B		Besson	7,653,031 7,672,714		3/2010	Godfrey et al.
7,218,967 B 7,231,451 B		Bergelson Law	7,673,679	B2	3/2010	Harrison et al.
7,243,118 B	2 7/2007	Lou	7,678,043 7,686,839		3/2010 3/2010	
7,246,521 B 7,249,212 B			7,697,994			VanDanacker et al.
7,252,792 B	2 8/2007	Perrault	7,720,036		5/2010	
7,253,716 B 7,261,690 B		Lovoi et al.	7,729,776 7,733,224		6/2010	Von Arx et al. Tran
7,270,633 B	9/2007	Goscha	7,736,318	B2	6/2010	Costentino
7,273,454 B	9/2007	Raymond et al.	7,756,587 7,760,104		7/2010 7/2010	Penner et al.
7,289,855 B 7,291,497 B		Nghiem Holmes	7,782,991	B2		Sobchak et al.
7,292,139 B	2 11/2007	Mazar et al.	7,796,043			Euliano et al.
7,294,105 B 7,311,665 B		Islam Hawthorne	7,797,033 7,809,399		10/2010	D'Andrea et al. Lu
7,313,163 B	2 12/2007		7,844,341	B2	11/2010	Von Arx et al.
7,317,378 B	2 1/2008	Jarvis et al.	7,881,799 7,975,587			Greenberg et al. Schneider
7,318,808 B 7,336,929 B	52 1/2008 52 2/2008	Tarassenko et al. Yasuda	7,978,064	B2	7/2011	Zdeblick et al.
7,342,895 B	2 3/2008	Serpa	7,983,189 8,036,731			Bugenhagen Kimchy et al.
7,346,380 B 7,349,722 B		Axelgaard et al. Witkowski et al.	8,036,731			Zdeblick et al.
7,352,998 B			8,054,047	B2		Chen et al.
7,353,258 B 7,357,891 B		Washburn Yang et al.	8,054,140 8,055,334			Fleming et al. Savage et al.
7,359,674 B		Markki	8,082,919	B2	12/2011	Brunnberg et al.
7,366,558 B		Virtanen et al.	8,119,045 8,131,376			Schmidt et al. Faraji et al.
7,368,190 B 7,368,191 B		Heller et al. Andelman et al.	8,177,611	B2	5/2012	Kang
7,373,196 B	5/2008	Ryu et al.	8,185,191			Shapiro et al. Headley
7,375,739 B 7,376,435 B		Robbins McGowan	8,185,646 8,200,320		6/2012	Kovacs
7,382,247 B	2 6/2008	Welch et al.	8,207,731		6/2012	Moskalenko
7,382,263 B 7,387,607 B		Danowski et al.	8,224,596 8,253,586		8/2012	Agrawal et al. Matak
7,388,903 B		Godfrey et al.	8,254,853	B2	8/2012	Rofougaran
7,389,088 B			8,271,146 8,298,574			Heber et al. Tsabari et al.
7,392,015 B 7,395,106 B	51 6/2008 52 7/2008	Farlow Ryu et al.	8,343,068	B2	1/2013	Najafi et al.
7,396,330 B	2 7/2008	Banet	8,374,698 8,389,003			Ok et al. Mintchev et al.
7,404,968 B 7,413,544 B		Abrams et al. Kerr	8,404,275		3/2013	Habboushe
7,414,534 B	1 8/2008	Kroll et al.	8,425,492 8,443,214			Herbert et al.
7,414,543 B 7,415,242 B		Rye et al.	8,443,214			Lee et al. Yuen et al.
7,424,268 B		Diener	8,532,776			Greenberg et al.
7,424,319 B 7,427,266 B		Muehlsteff Ayer et al.	8,540,664 8,545,402			Robertson et al. Hafezi et al.
7,442,164 B		Berrang et al.	8,547,248	B2	10/2013	Zdeblick et al.
7,443,290 B	2 10/2008	Takiguchi	8,558,563 8,564,432			Zdeblick Covannon et al.
7,458,887 B 7,469,838 B		Kurosawa Brooks et al.	8,597,186			Hafezi et al.
7,471,665 B	2 12/2008	Perlman	8,634,838		1/2014	Hellwig et al.
7,471,992 B 7,492,128 B		Schmidt et al.	8,647,358 8,660,645			Brister et al. Stevenson et al.
7,492,128 B 7,499,674 B		Salokannel	8,668,643		3/2014	
7,510,121 B	3/2009	Koenck	8,685,451		4/2014	Toneguzzo et al.
7,512,448 B 7,515,043 B		Malick Welch	8,697,057 8,698,006			Van Epps et al. Bealka et al.
7,513,043 B 7,519,416 B		Sula et al.	8,758,237	B2		Sherman et al.
7,523,756 B	2 4/2009	Minai	8,784,308		7/2014	Duck et al.

# **US 10,542,909 B2**Page 5

(56)	Referei	nces Cited	2003/0162556 A1	8/2003		
U.S.	PATENT	DOCUMENTS	2003/0164401 A1 2003/0167000 A1		Andreasson et al. Mullick et al.	
0.5.	11112111	Bocomercia	2003/0171791 A1		KenKnight	
8,802,183 B2		Frank et al.	2003/0171898 A1		Tarassenko et al. Yokoi et al.	
8,816,847 B2		Zdeblick et al.	2003/0181788 A1 2003/0185286 A1	10/2003		
8,836,513 B2 8,838,217 B2	9/2014	Hafezi et al. Myr	2003/0187337 A1	10/2003	Tarassenko et al.	
8,858,432 B2		Robertson	2003/0187338 A1		Say et al.	
8,908,943 B2		Berry et al.	2003/0195403 A1 2003/0213495 A1		Berner et al. Fuita et al.	
8,912,908 B2		Berkman et al.	2003/0213495 A1 2003/0214579 A1	11/2003	- · J	
8,926,509 B2 8,932,221 B2		Magar et al. Colliou et al.			Bunick A61K 9/2	866
8,945,005 B2		Hafezi et al.			427	/2.1
8,951,234 B2		Hafezi et al.	2003/0216622 A1		Meron et al.	
8,989,837 B2 9,031,658 B2		Weinstein et al. Chiao et al.	2003/0216625 A1 2003/0216666 A1	11/2003	Ericson et al.	
9,088,168 B2		Mach et al.	2003/0216729 A1		Marchitto	
9,107,806 B2	8/2015	Hafezi et al.	2003/0219484 A1		Sowden et al.	
9,119,554 B2		Robertson et al.	2003/0232895 A1 2004/0008123 A1		Omidian et al. Carrender et al.	
9,119,918 B2 9,149,423 B2		Robertson et al. Duck et al.	2004/0008123 A1 2004/0018476 A1		LaDue	
9,158,890 B2		Meredith et al.	2004/0034295 A1		Salganicoff	
9,161,707 B2		Hafezi et al.	2004/0049245 A1	3/2004		
9,189,941 B2 9,226,663 B2	11/2015 1/2016	Eschelman et al.	2004/0073095 A1 2004/0073454 A1		Causey et al. Urquhart et al.	
9,226,6679 B2		Balda	2004/0077995 A1		Ferek-Petric	
9,268,909 B2		Jani et al.	2004/0082982 A1		Gord et al.	
9,270,025 B2		Robertson et al.	2004/0087839 A1 2004/0092801 A1		Raymond et al. Drakulic	
9,271,897 B2 9,277,864 B2		Costello et al. Yang et al.	2004/0092801 A1 2004/0106859 A1		Say et al.	
9,320,455 B2		Hafezi et al.	2004/0115507 A1	6/2004	Potter et al.	
9,415,010 B2		Hafezi et al.	2004/0115517 A1		Fukada et al.	
9,433,371 B2		Hafezi et al. Berkman et al.	2004/0117062 A1 2004/0121015 A1		Bonney Chidlaw et al.	
9,439,582 B2 9,439,599 B2		Thompson et al.	2004/0148140 A1		Tarassenko et al.	
9,517,012 B2	12/2016	Lane et al.	2004/0153007 A1	8/2004		
9,597,010 B2		Thompson et al.	2004/0167226 A1 2004/0167465 A1		Serafini Mihai et al.	
9,597,487 B2 9,599,679 B2		Robertson et al. Taylor et al.	2004/0167801 A1		Say et al.	
9,649,066 B2		Zdeblick et al.	2004/0193020 A1	9/2004	Chiba	
9,681,842 B2		Zdeblick et al.	2004/0193029 A1		Gluhovsky Mayer et al.	
9,741,975 B2 9,756,874 B2		Laulicht et al. Arne et al.	2004/0193446 A1 2004/0199222 A1		Sun et al.	
9,968,284 B2		Vidalis et al.	2004/0215084 A1		Shimizu et al.	
2001/0027331 A1		Thompson	2004/0218683 A1	11/2004		
2001/0044588 A1 2001/0051766 A1	11/2001	Mault Gazdinski	2004/0220643 A1 2004/0224644 A1	11/2004	Schmidt Wu	
2001/0031/00 A1 2002/0002326 A1		Causey et al.	2004/0225199 A1	11/2004		
2002/0026111 A1	2/2002	Ackerman	2004/0253304 A1*	12/2004	Gross A61B 1/	
2002/0032384 A1		Raymond et al. Raymond et al.	2004/0258571 A1	12/2004	Lee et al.	451
2002/0032385 A1 2002/0040278 A1		Anuzis et al.			Sanghvi A61K 9/0	031
2002/0077620 A1		Sweeney et al.			514/	
2002/0099423 A1		Berg et al.	2004/0260154 A1		Sidelnik	
2002/0128934 A1 2002/0132226 A1	9/2002	Shaer Nair	2005/0003074 A1 2005/0017841 A1	1/2005	Brown et al.	
2002/0136744 A1		McGlynn et al.	2005/0020887 A1		Goldberg	
2002/0179921 A1	12/2002		2005/0021370 A1	1/2005		
2002/0192159 A1 2002/0193669 A1		Reitberg Glukhovsky	2005/0024198 A1 2005/0027205 A1	2/2005	Ward Tarassenko et al.	
2002/0193009 A1 2002/0198470 A1	12/2002	Imran et al.	2005/0027203 A1 2005/0038321 A1		Fujita et al.	
2003/0017826 A1	1/2003	Fishman et al.	2005/0043634 A1	2/2005	Yokoi et al.	
2003/0023150 A1		Yokoi et al.	2005/0043894 A1		Fernandez	
2003/0028226 A1 2003/0039956 A1*		Thompson Choi A61K 9/1652	2005/0054897 A1 2005/0055014 A1		Hashimoto et al. Coppeta et al.	
		435/5	2005/0062644 A1	3/2005	Leci	
2003/0062551 A1		Chen et al.	2005/0065407 A1		Nakamura et al.	
2003/0065536 A1 2003/0076179 A1		Hansen Branch et al.	2005/0070778 A1 2005/0075145 A1		Lackey Dvorak et al.	
2003/00/61/9 A1 2003/0083559 A1		Thompson	2005/0090753 A1		Goor et al.	
2003/0091625 A1	5/2003	Hariharan et al.	2005/0092108 A1		Andermo	
2003/0126593 A1		Mault Nielsen et al.	2005/0096514 A1 2005/0096562 A1		Starkebaum Delalic et al.	
2003/0130714 A1 2003/0135128 A1		Suffin et al.	2005/0090302 AT 2005/0101843 AT	5/2005		
2003/0135392 A1	7/2003	Vrijens et al.	2005/0101872 A1		Sattler	
2003/0152622 A1		Louie-Helm et al.	2005/0115561 A1		Stahmann et al.	
2003/0158466 A1 2003/0158756 A1		Lynn et al. Abramson	2005/0116820 A1 2005/0117389 A1		Goldreich Worledge	
2005/0150/30 Al	0/2003	1 WIAIIISUII	2003/011/3 <b>6</b> 9/Al	0/2003	wonedge	

# US 10,542,909 B2 Page 6

(56) R	References Cited	2007/0006636 A1	1/2007 King et al.
,	TENET DOCLINAENTS	2007/0008113 A1	1/2007 Spoonhower et al. 1/2007 Fischell et al.
U.S. PA	TENT DOCUMENTS	2007/0016089 A1 2007/0027386 A1	2/2007 Fischen et al.
2005/0121322 A1	6/2005 Say et al.	2007/0027388 A1	2/2007 Chou
	6/2005 Ayer et al.	2007/0029195 A1	2/2007 Li et al.
	6/2005 Kojima	2007/0038054 A1	2/2007 Zhou
	7/2005 Nakatani et al.	2007/0049339 A1 2007/0055098 A1	3/2007 Barak et al. 3/2007 Shimizu et al.
	7/2005 Boesen 7/2005 Bruinsma	2007/0060797 A1	3/2007 Ball
	7/2005 Gilbert et al.	2007/0060800 A1	3/2007 Drinan et al.
	7/2005 Montgomery	2007/0066929 A1	3/2007 Ferren et al.
	8/2005 Takizawa	2007/0073353 A1 2007/0096765 A1	3/2007 Rooney et al. 5/2007 Kagan
	8/2005 LaPorte 8/2005 Hatlestad et al.	2007/0106346 A1	5/2007 Bergelson
	9/2005 Marshall	2007/0123772 A1	5/2007 Euliano
	9/2005 DelMain et al.	2007/0129622 A1	6/2007 Bourget
	9/2005 Aisenbrey	2007/0130287 A1 2007/0135803 A1	6/2007 Kumar 6/2007 Belson
	0/2005 Cole 0/2005 Heinonen	2007/0142721 A1	6/2007 Berner et al.
	0/2005 Remoden 0/2005 Bogash et al.	2007/0156016 A1	7/2007 Betesh
2005/0245794 A1 1	1/2005 Dinsmoor	2007/0160789 A1 2007/0162089 A1	7/2007 Merical
	1/2005 Yang et al.	2007/0162089 A1 2007/0162090 A1	7/2007 Mosesov 7/2007 Penner
	1/2005 Mumford 2/2005 Shuros et al.	2007/0167495 A1	7/2007 Brown et al.
	2/2005 Schultz et al.	2007/0167848 A1	7/2007 Kuo et al.
2005/0277912 A1 1:	2/2005 John	2007/0173701 A1	7/2007 Al-Ali
	2/2005 Strother et al.	2007/0179347 A1 2007/0179371 A1	8/2007 Tarassenko et al. 8/2007 Peyser et al.
	2/2005 Mauze et al. 2/2005 Pettus	2007/0185393 A1	8/2007 Zhou
	2/2005 Tettus 2/2005 Sengupta	2007/0191002 A1	8/2007 Ge
2005/0288594 A1 11	2/2005 Lewkowicz et al.	2007/0196456 A1	8/2007 Stevens
	1/2006 Abrosimov et al.	2007/0207793 A1 2007/0208233 A1	9/2007 Myer 9/2007 Kovacs
	2/2006 Moon et al. 2/2006 Tarassenko et al.	2007/0213659 A1	9/2007 Trovato et al.
	3/2006 Kwiatkowski et al.	2007/0237719 A1	10/2007 Jones
	3/2006 Lovoi et al.	2007/0244370 A1	10/2007 Kuo et al.
	3/2006 Kingery	2007/0255198 A1 2007/0255330 A1	11/2007 Leong et al. 11/2007 Lee
	3/2006 Begleiter 4/2006 Henderson	2007/0270672 A1	11/2007 Lee 11/2007 Hayter
	4/2006 Barnes et al.	2007/0279217 A1	12/2007 Venkatraman
	4/2006 Yang et al.	2007/0282174 A1	12/2007 Sabatino
	5/2006 Drew	2007/0282177 A1 2007/0299480 A1	12/2007 Pilz 12/2007 Hill
	5/2006 Bettesh et al. 5/2006 Han	2008/0000804 A1	1/2008 Carey et al.
	5/2006 Keating	2008/0014866 A1	1/2008 Lipowshi
	5/2006 Powell	2008/0020037 A1 2008/0021519 A1	1/2008 Robertson et al. 1/2008 DeGeest
	6/2006 Teller et al. 6/2006 Bouchoucha	2008/0021519 A1 2008/0021521 A1	1/2008 Decleest 1/2008 Shah
	6/2006 Chavan et al.	2008/0027679 A1	1/2008 Shklarski
	6/2006 Lee et al.	2008/0033273 A1	2/2008 Zhou
	6/2006 Tarassenko et al.	2008/0038588 A1 2008/0039700 A1	2/2008 Lee 2/2008 Drinan et al.
	6/2006 Banet 7/2006 Kimura	2008/0035700 A1 2008/0045843 A1	2/2008 Tsuji et al.
	7/2006 McLean	2008/0046038 A1	2/2008 Hill
	7/2006 Burnes	2008/0051647 A1	2/2008 Wu et al.
	7/2006 Glukhovsky et al. 7/2006 Kroecker	2008/0051667 A1 2008/0058614 A1	2/2008 Goldreich 3/2008 Banet
	7/2006 Kroecker 7/2006 Sormann et al.	2008/0062856 A1	3/2008 Feher
	8/2006 Kim	2008/0065168 A1	3/2008 Bitton et al.
	8/2006 Horn	2008/0074307 A1 2008/0077015 A1	3/2008 Boric-Lubecke 3/2008 Botic-Lubecke
	8/2006 Atanasoska et al. 9/2006 Dubrow et al.	2008/0077013 A1 2008/0077028 A1	3/2008 Schaldach et al.
	9/2006 Dublow et al. 9/2006 Spaeder	2008/0077188 A1	3/2008 Denker et al.
	9/2006 Choi	2008/0091089 A1	4/2008 Guillory et al.
	9/2006 Walker	2008/0091114 A1 2008/0097549 A1	4/2008 Min 4/2008 Colbaugh
	0/2006 Drew 1/2006 Kim et al.	2008/0097917 A1	4/2008 Colbaugh 4/2008 Dicks
	1/2006 Kim et al. 1/2006 Siddiqui	2008/0103440 A1	5/2008 Ferren et al.
2006/0253005 A1 1	1/2006 Drinan	2008/0112885 A1	5/2008 Okunev et al.
	1/2006 Ibrahim	2008/0114224 A1	5/2008 Bandy et al.
	2/2006 Posamentier 2/2006 McGinnis	2008/0119705 A1 2008/0119716 A1	5/2008 Patel 5/2008 Boric-Lubecke
	2/2006 McGrinns 2/2006 Pinkney	2008/0119710 A1 2008/0121825 A1	5/2008 Trovato et al.
	2/2006 Noel	2008/0137566 A1	6/2008 Marholev
	2/2006 Mercure	2008/0139907 A1	6/2008 Rao et al.
	2/2006 Alt	2008/0140403 A1	6/2008 Hughes et al.
	1/2007 Karube et al. 1/2007 Suzuki	2008/0146871 A1 2008/0146889 A1	6/2008 Arneson et al. 6/2008 Young
200 11 000 2030 A1	1/2007 Suzuki	2000/0170007 A1	orzood Toung

# **US 10,542,909 B2**Page 7

(56)	References Cited	2009/0177060 A1	7/2009	
IIC	DATENT DOCUMENTS	2009/0177061 A1 2009/0177062 A1	7/2009 7/2009	Say et al. Say et al.
0.5.	PATENT DOCUMENTS	2009/0177062 A1 2009/0177063 A1	7/2009	Say et al.
2008/0146892 A1	6/2008 LeBeouf	2009/0177064 A1	7/2009	Say et al.
2008/0154104 A1	6/2008 Lamego	2009/0177065 A1	7/2009	Say et al.
2008/0166992 A1	7/2008 Ricordi	2009/0177066 A1	7/2009	•
2008/0175898 A1	7/2008 Jones et al.	2009/0182206 A1 2009/0182207 A1	7/2009 7/2009	
2008/0183245 A1	7/2008 Van Oort 8/2008 Belsky et al.	2009/0182207 A1 2009/0182212 A1	7/2009	Say et al.
2008/0188837 A1 2008/0194912 A1	8/2008 Trovato et al.	2009/0182213 A1	7/2009	
2008/0208009 A1	8/2008 Shklarski	2009/0182214 A1	7/2009	
2008/0214901 A1	9/2008 Gehman	2009/0182215 A1	7/2009	
2008/0214985 A1	9/2008 Yanaki	2009/0182388 A1 2009/0187088 A1	7/2009 7/2009	Von Arx Say et al.
2008/0243020 A1 2008/0249360 A1	10/2008 Chou 10/2008 Li	2009/0187089 A1	7/2009	Say et al.
2008/0262320 A1	10/2008 Schaefer et al.	2009/0187090 A1	7/2009	Say et al.
2008/0262336 A1	10/2008 Ryu	2009/0187091 A1	7/2009	
2008/0269664 A1	10/2008 Trovato et al.	2009/0187092 A1 2009/0187093 A1	7/2009 7/2009	
2008/0275312 A1 2008/0284599 A1*	11/2008 Mosesov * 11/2008 Zdeblick et al A61B 5/0031	2009/0187094 A1	7/2009	•
2000/020 <del>1</del> 333 A1	340/572.1	2009/0187095 A1	7/2009	
2008/0288027 A1	11/2008 Kroll	2009/0187381 A1	7/2009	
2008/0294020 A1	11/2008 Sapounas	2009/0192351 A1 2009/0192368 A1	7/2009 7/2009	
2008/0299197 A1	12/2008 Toneguzzo et al.	2009/0192369 A1	7/2009	Say et al.
2008/0300572 A1 2008/0303638 A1	12/2008 Rankers 12/2008 Nguyen	2009/0192370 A1	7/2009	Say et al.
2008/0306357 A1	12/2008 Korman	2009/0192371 A1	7/2009	Say et al.
2008/0306359 A1	12/2008 Zdeblick et al.	2009/0192372 A1	7/2009	
2008/0306360 A1	12/2008 Robertson et al.	2009/0192373 A1 2009/0192374 A1	7/2009 7/2009	
2008/0311852 A1 2008/0312522 A1	12/2008 Hansen 12/2008 Rowlandson	2009/0192375 A1	7/2009	
2008/0312322 A1 2008/0316020 A1	12/2008 Robertson	2009/0192376 A1	7/2009	Say et al.
2009/0009330 A1	1/2009 Sakama et al.	2009/0192377 A1	7/2009	•
2009/0009332 A1	1/2009 Nunez et al.	2009/0192378 A1 2009/0192379 A1	7/2009 7/2009	
2009/0024045 A1	1/2009 Prakash	2009/0192377 A1 2009/0194747 A1	8/2009	,
2009/0024112 A1 2009/0030293 A1	1/2009 Edwards et al. 1/2009 Cooper et al.	2009/0197068 A1	8/2009	Yamaguchi et al.
2009/0030297 A1	1/2009 Miller	2009/0198115 A1	8/2009	Say et al.
2009/0034209 A1	2/2009 Joo	2009/0198116 A1 2009/0198175 A1	8/2009 8/2009	•
2009/0043171 A1	2/2009 Rule	2009/0198173 A1 2009/0203964 A1	8/2009	
2009/0047357 A1 2009/0048498 A1	2/2009 Tomohira et al. 2/2009 Riskey	2009/0203971 A1	8/2009	Sciarappa
2009/0062634 A1	3/2009 Say et al.	2009/0203972 A1	8/2009	
2009/0062670 A1	3/2009 Sterling	2009/0203978 A1 2009/0204265 A1	8/2009 8/2009	Say et al. Hackett
2009/0062730 A1	3/2009 Woo	2009/0204203 A1 2009/0210164 A1	8/2009	
2009/0069642 A1 2009/0069655 A1	3/2009 Gao 3/2009 Say et al.	2009/0216101 A1	8/2009	Say et al.
2009/0069656 A1	3/2009 Say et al.	2009/0216102 A1	8/2009	Say et al.
2009/0069657 A1	3/2009 Say et al.	2009/0227204 A1 2009/0227876 A1	9/2009 9/2009	Robertson et al.
2009/0069658 A1	3/2009 Say et al.	2009/0227940 A1		Say et al.
2009/0069724 A1 2009/0076343 A1	3/2009 Otto et al. 3/2009 James	2009/0227941 A1	9/2009	
2009/0076350 A1	3/2009 Bly et al.	2009/0227988 A1	9/2009	
2009/0082645 A1	3/2009 Hafezi et al.	2009/0228214 A1 2009/0231125 A1	9/2009 9/2009	
2009/0087483 A1 2009/0088618 A1	4/2009 Sison	2009/0234200 A1		Husheer
2009/0088618 A1 2009/0099435 A1	4/2009 Ameson 4/2009 Say et al.	2009/0243833 A1	10/2009	
2009/0105561 A1	4/2009 Boydon et al.	2009/0253960 A1	10/2009	Takenaka et al.
2009/0110148 A1	4/2009 Zhang	2009/0256702 A1 2009/0260212 A1	10/2009 10/2009	Robertson Schmett et al.
2009/0112626 A1	4/2009 Talbot	2009/0264714 A1	10/2009	
2009/0124871 A1 2009/0124965 A1	5/2009 Arshak 5/2009 Greenberg et al.	2009/0264964 A1	10/2009	Abrahamson
2009/0131774 A1	5/2009 Sweitzer	2009/0265186 A1	10/2009	
2009/0135886 A1	5/2009 Robertson et al.	2009/0273467 A1 2009/0281539 A1	11/2009	Elixmann Selig
2009/0142853 A1	6/2009 Warrington et al.	2009/0281339 A1 2009/0287109 A1		Ferren et al.
2009/0149839 A1 2009/0157113 A1	6/2009 Hyde et al. 6/2009 Marcotte	2009/0295548 A1		Ronkka
2009/0157115 A1 2009/0157358 A1	6/2009 Kim	2009/0296677 A1	12/2009	•
2009/0161602 A1	6/2009 Matsumoto	2009/0303920 A1	12/2009	
2009/0163789 A1	6/2009 Say et al.	2009/0306633 A1 2009/0312619 A1	12/2009 12/2009	
2009/0171180 A1 2009/0171420 A1	7/2009 Pering 7/2009 Brown et al.	2009/0312019 A1 2009/0318303 A1	12/2009	•
2009/0171420 A1 2009/0173628 A1	7/2009 Say et al.	2009/0318761 A1		Rabinovitz
2009/0177055 A1	7/2009 Say et al.	2009/0318779 A1	12/2009	Tran
2009/0177056 A1	7/2009 Say et al.	2009/0318783 A1	12/2009	
2009/0177057 A1	7/2009 Say et al.	2009/0318793 A1	1/2009	
2009/0177058 A1 2009/0177059 A1	7/2009 Say et al. 7/2009 Say et al.	2010/0001841 A1 2010/0010330 A1		Cardullo Rankers
2009/01/7039 AI	112009 Say St al.	2010/0010330 Al	1/2010	Name 13

## US 10,542,909 B2

Page 8

(56)	Referer	ices Cited			31283 A1		Wang et al.
U.S.	PATENT	DOCUMENTS		2013/01	71596 A1 72690 A1	7/2013	French Arne et al.
					85228 A1		Dresner
2010/0033324 A1		Shimizu et al.			96012 A1 99662 A1	8/2013	Dill Gebbink
2010/0036269 A1 2010/0049004 A1		Ferren et al. Edman et al.			09877 A1		Kren et al.
2010/0049004 A1 2010/0049006 A1		Magar			23028 A1	8/2013	Arne et al.
2010/0049012 A1		Dijksman et al.			75296 A1		Tietzen et al.
2010/0049069 A1		Tarassenko et al.			66734 A1 79221 A1		Zdeblick Whitworth et al.
2010/0056878 A1 2010/0056891 A1		Partin Say et al.			80202 A1		Zdeblick et al.
2010/0056939 A1		Tarassenko et al.			80125 A1		Bhardwaj et al.
2010/0057041 A1		Hayter			08930 A1	10/2014	
2010/0062709 A1	3/2010				49256 A1 74276 A1	11/2014	Connor Guthrie et al.
2010/0063438 A1 2010/0063841 A1		Bengtsson D'Ambrosia et al.			17486 A1	1/2015	
2010/0003841 A1 2010/0069002 A1	3/2010				59922 A1		Thompson et al.
2010/0069717 A1		Hafezi et al.			80678 A1		Frank et al.
2010/0099967 A1		Say et al.			80679 A1 80680 A1		Frank et al. Zdeblick et al.
2010/0099968 A1 2010/0099969 A1		Say et al. Say et al.			12243 A1		Hafezi et al.
2010/0100077 A1	4/2010				27737 A1	5/2015	
2010/0100078 A1	4/2010	Say et al.			27738 A1		Thompson et al.
2010/0106001 A1		Say et al.			49375 A1 50480 A1		Thompson et al. Zdeblick et al.
2010/0118853 A1 2010/0139672 A1		Godfrey Kroll et al.			64746 A1		Costello et al.
2010/0168659 A1		Say et al.			30729 A1		Zdeblick et al.
2010/0179398 A1		Say et al.			48833 A1 52343 A1		Arne et al.
2010/0191073 A1		Tarassenko et al. Gorbachov			61234 A1		Hafezi et al. Hafezi et al.
2010/0210299 A1 2010/0222652 A1	9/2010				33667 A1		Schmidt et al.
2010/0228113 A1		Solosko			45906 A1		Johnson et al.
2010/0233026 A1		Ismagliov et al.			80708 A1 00179 A1		Dua et al. Cheng et al.
2010/0234706 A1 2010/0234715 A1	9/2010 9/2010	Gilland			14046 A1		Hafezi et al.
2010/0234713 A1 2010/0234914 A1	9/2010				20182 A1		Schmidt et al.
2010/0239616 A1*		Hafezi	A61B 5/073		16569 A1		Hafezi et al.
2010/02/2001	0.0040	at t	424/400		65813 A1 74194 A1		Zdeblick et al. Robertson et al.
2010/0245091 A1 2010/0249541 A1	9/2010	Singh Geva et al.			96799 A1		Hafezi et al.
2010/0249341 A1 2010/0249881 A1		Corndorf		2018/00	26680 A1		Shirvani et al.
2010/0256461 A1		Mohamedali			10441 A1		Frank et al.
2010/0259543 A1		Tarassenko et al.			84698 A1 14048 A1		Arne et al. Zdeblick et al.
2010/0268048 A1 2010/0268049 A1		Say et al. Say et al.			29996 A1		Thompson
2010/0268050 A1	10/2010	Say et al.					1
2010/0274111 A1		Say et al.			FOREIG	GN PATE	NT DOCUMENTS
2010/0280345 A1 2010/0280346 A1		Say et al. Say et al.		CDI	1015	5202	0/2010
2010/0280540 A1 2010/0295694 A1		Kauffman et al.		CN DE	10179	)5202 13005	8/2010 10/2004
2010/0297640 A1	11/2010	Kumar et al.		EP		14939	12/1989
2010/0298650 A1		Moon et al.		EP	052	26166	2/1993
2010/0298668 A1 2010/0298730 A1		Hafezi et al. Tarassenko et al.		EP EP		31152	2/2000
2010/0312188 A1		Robertson et al.		EP EP		16356 34054	10/2002 5/2005
2010/0312580 A1		Tarassenko et al.		EP	170	)2553	9/2006
2011/0009715 A1 2011/0065983 A1		O'Reilly et al. Hafezi et al.		EP		14308	12/2007
2011/0077660 A1		Janik et al.		EP GB		13369 27762	1/2010 2/1960
2011/0124983 A1		Kroll et al.		JP		2712	4/1986
2011/0134906 A1 2011/0160549 A1		Garudadri et al. Saroka et al.		JP	H0128		11/1989
2011/0100349 A1 2011/0224912 A1		Bhavaraju et al.		JP JP	0522 H1119	28128	9/1993 7/1999
2011/0230732 A1		Edman et al.		JР	200050		5/2000
2011/0270135 A1		Dooley et al.		JP	200226		9/2002
2012/0004520 A1 2012/0011699 A1		Whitworth et al. Hafezi et al.		JP	200228		10/2002
2012/0011033 A1 2012/0016231 A1		Westmoreland		JP JP	200305 200431		2/2003 11/2004
2012/0032816 A1	2/2012	Cho et al.		JР	200431		3/2005
2012/0062371 A1		Radivojevic et al.		JP	200508	37552	4/2005
2012/0071743 A1 2012/0109112 A1	3/2012 5/2012	Todorov et al. Strand et al.		JP ID	200510		4/2005 5/2005
2012/0179004 A1	7/2012	Roesicke et al.		JP JP	200512 200551		5/2005 5/2005
2012/0245043 A1		England		JP	200530		11/2005
2012/0276451 A1		Lestriez et al.		JР	200534		12/2005
2012/0299723 A1 2013/0129869 A1		Hafezi et al. Hafezi et al.		JP JP	2005533 200600		12/2005 1/2006
2013/0129809 A1 2013/0129872 A1		Kruger		JР	200650		3/2006
	_	-					

(56)	References Cited	WO WO2007071180 6/2007
()		WO WO2007096810 8/2007 TS WO WO2007101141 9/2007
	FOREIGN PATENT DOCUMEN	TS WO WO2007101141 9/2007 WO WO2007115087 10/2007
JР	2007200739 8/2007	WO WO2007120946 10/2007
JP	2007313340 12/2007	WO WO2007127316 11/2007 WO WO2007127879 11/2007
JP JP	2009514870 4/2009	WO WO2007127879 11/2007 WO WO2007128165 11/2007
KR	2009528909 8/2009 200600977523 7/2006	WO WO2007130491 11/2007
TW	200406192 5/2004	WO WO2007143535 12/2007
TW	200916136 4/2009	WO WO2007149546 12/2007 WO WO2008008281 1/2008
WO WO	WO1988002237 4/1988 WO1992021307 12/1992	WO WO2008012700 1/2008
wo	WO1993008734 5/1993	WO WO2008030482 3/2008
WO	WO1993019667 10/1993	WO WO2008052136 5/2008 WO WO-2008052136 A2 * 5/2008
WO WO	WO1994001165 1/1994 WO1997039963 10/1997	WO WO2008063626 5/2008
WO	WO1998043537 10/1998	WO WO2008066617 6/2008
WO	WO1999037290 7/1999	WO WO2008076464 6/2008 WO WO2008089232 7/2008
WO WO	WO1999059465 11/1999 WO2000032474 6/2000	WO WO2008089232 7/2008 WO WO2008091683 7/2008
WO	WO2000032474 6/2000 WO2000033246 6/2000	WO WO2008095183 8/2008
WO	WO2001000085 1/2001	WO WO2008097652 8/2008
WO	WO2001047466 7/2001	WO WO2008101107 8/2008 WO WO2008112577 9/2008
WO WO	WO2001058236 8/2001 WO2001074011 10/2001	WO WO2008112578 9/2008
WO	WO2001080731 11/2001	WO WO2008120156 10/2008
WO	WO2002000920 1/2002	WO WO2008133394 11/2008 WO WO2008134185 11/2008
WO WO	WO2002045489 6/2002 WO2002058330 7/2002	WO WO2008154163 11/2008 WO WO2008150633 12/2008
WO	WO2002038330 7/2002 WO2002062276 8/2002	WO WO2009000447 12/2008
WO	WO2002087681 11/2002	WO WO2009001108 12/2008
WO WO	WO2002095351 11/2002 WO2003005877 1/2003	WO WO2009006615 1/2009 WO WO2009029453 3/2009
WO	WO2003003877 1/2003 WO2003050643 6/2003	WO WO2009031149 3/2009
WO	WO2003068061 8/2003	WO WO2009036334 3/2009
WO	WO2004014225 2/2004	WO WO2009051829 4/2009 WO WO2009051830 4/2009
WO WO	WO2004019172 3/2004 WO2004039256 5/2004	WO WO2009063377 5/2009
WO	WO2004066833 8/2004	WO WO2009081348 7/2009
WO	WO2004066834 8/2004	WO WO2009111664 9/2009 WO WO2009146082 12/2009
WO WO	WO2004066903 8/2004 WO2004068881 8/2004	WO WO2009140082 12/2009 WO WO2010009100 1/2010
WO	WO2004003831 8/2004 WO2004075032 9/2004	WO WO2010011833 1/2010
WO	WO2004109316 12/2004	WO WO2010019778 2/2010 WO WO2010057049 5/2010
WO WO	WO2005011237 2/2005 WO2005020023 3/2005	WO WO2010037049 3/2010 WO WO2010080765 7/2010
WO	WO2005024687 3/2005	WO WO2010080843 7/2010
WO	WO2005041438 5/2005	WO WO2010107563 9/2010 WO WO2010129288 11/2010
WO WO	WO2005047837 5/2005 WO2005051166 6/2005	WO WO2010129288 11/2010 WO WO2010132331 11/2010
WO	WO20050531100 0/2005 WO2005053517 6/2005	WO WO2010135516 11/2010
WO	WO2005083621 9/2005	WO WO2011068963 6/2011
WO WO	WO2005110238 11/2005 WO2005123569 12/2005	WO WO2011133799 10/2011 WO WO2011159336 12/2011
WO	WO2005123569 12/2005 WO2006021932 3/2006	WO WO2011159337 12/2011
WO	WO2006027586 3/2006	WO WO2011159338 12/2011
WO	WO2006028347 3/2006	WO WO2011159339 12/2011 WO WO2012112561 8/2012
WO WO	WO2006055892 5/2006 WO2006055956 5/2006	WO WO2015112603 7/2015
WO	WO2006075016 7/2006	WO WO2015112604 7/2015
WO	WO2006100620 9/2006	WO WO2015119911 8/2015
WO WO	WO2006104843 10/2006 WO2006116718 11/2006	OTHER RIDITORIC
WO	WO-2006116718 A2 * 11/2006	OTHER PUBLICATIONS
WO	WO2006127355 11/2006	Arshak et al., A Review and Adaptation of Methods of Object
WO WO	WO2007001724 1/2007 WO2007001742 1/2007	Tracking to Telemetry Capsules IC-Med (2007) vol. 1, No. 1, Issue
WO	WO2007001742 1/2007 WO2007013952 2/2007	1, 12pp.
WO	WO2007014084 2/2007	"ASGE Technology Status Evaluation Report: wireless capsule
WO WO	WO2007014527 2/2007 WO2007021496 2/2007	endoscopy" American Soc. for Gastrointestinal Endoscopy (2006) vol. 63, No. 4; 7 pp.
WO	WO2007021496 2/2007 WO2007027660 3/2007	Au-Yeung, K., et al., "A Networked System for Self-Management
WO	WO2007028035 3/2007	of Drug Therapy and Wellness", Wireless Health '10, Oct. 5-7,
WO	WO2007036687 4/2007	2010, San Diego, 9 pp.
WO	WO2007036741 4/2007	Aydin et al., "Design and implementation considerations for an
WO WO	WO2007036746 4/2007 WO2007040878 4/2007	advanced wireless interface in miniaturized integrated sensor Microsystems" Sch. of Eng. & Electron., Edinburgh Univ., UK;
WO	WO2007067054 6/2007	(2003); abstract (1 page).
		V <b>x</b> ♥ /

### (56) References Cited

### OTHER PUBLICATIONS

Barrie, Heidelberg pH capsule gastric analysis. Texbook of Natural Medicine, (1992), Pizzorno, Murray & Barrie (4 pages).

Bohidar et al., "Dielectric Behavior of Gelatin Solutions and Gels" Colloid Polym Sci (1998) 276:81-86.

Brock, "Smart Medicine: The Application of Auto-ID Technology to Healthcare" Auto-ID Labs (2002) http://www.autoidlabs.org/uploads/media/MIT-AUTOID-WH-010.pdf (14 pages).

Carlson et al., "Evaluation of a non-invasive respiratory monitoring system for sleeping subjects" Physiological Measurement (1999) 20(1): 53

Coury, L. "Conductance Measurement Part 1: Theory"; Current Separations, 18:3 (1999) p. 91-96.

Delvaux et al., "Capsule endoscopy: Technique and indications" Clinical Gastroenterology (2008) vol. 22, Issue 5, pp. 813-837.

Description of ePatch Technology Platform for ECG and EMG, located it http://www.madebydelta.com/imported/images/DELTA\_Web/documents/ME/ePatch\_ECG\_EMG.pdf, Dated Sep. 2, 2010; 1 page.

Dhar et al., "Electroless nickel plated contacts on porous silicon" Appl. Phys. Lett. 68 (10) pp. 1392-1393 (1996).

Eldek A., "Design of double dipole antenna with enhanced usable bandwidth for wideband phased array applications" Progress in Electromagnetics Research PIER 59, 1-15 (2006).

Fawaz et al., "Enhanced Telemetry System using CP-QPSK Band—Pass Modulation Technique Suitable for Smart Pill Medical Application" IFIP IEEE Dubai Conference (2008); http://www.asic.fh-offenburg.de/downloads/ePille/IFIP\_IEEE\_Dubai\_Conference.pdf (5 pages).

Ferguson et al., "Dielectric Constant Studies III Aqueous Gelatin Solutions" J. Chem. Phys. 2, 94 (1934) p. 94-98.

Furse C. M., "Dipole Antennas" J. Webster (ed). Wiley Encyclopedia of Electrical and Electronics Engineering (1999) p. 575-581. Gaglani S. "Put Your Phone, or Skin, on Vibrate" MedGadget (2012) http://medgadget.com/2012/03/put-your-phone-or-skin-on-vibrate.html 8pp.

Gilson, D.R. "Molecular dynamics simulation of dipole interactions", Department of Physics, Hull University, Dec. (2002), p. 1-43.

Given Imaging, "Agile Patency Brochure" (2006) http://www.inclino.no/documents/AgilePatencyBrochure\_Global\_GMB-0118-01.pdf; 4pp. Gonzalez-Guillaumin et al., "Ingestible capsule for impedance and pH monitoring in the esophagus" IEEE Trans Biomed Eng. (2007) 54(12): 2231-6; abstract.

Greene, "Edible RFID microchip monitor can tell if you take your medicine" Bloomberg Businessweek (2010) 2 pp.; http://www.businessweek.com/idg/2010-03-31/edible-rfid-microchip-monitor-can-tell-if-you-take-your-medicine.html (1 page).

Heydari et al., "Analysis of the PLL jitter due to power/ground and substrate noise"; IEEE Transactions on Circuits and Systems (2004) 51(12): 2404-16.

Hoeksma, J. "New 'smart pill' to track adherence" E-Health-Insider (2010) http://www.e-health-insider.com/news/5910/new\_'smart\_pill'\_monitors\_medicines (1 page).

Hoover et al., "Rx for health: Engineers design pill that signals it has been swallowed" University of Florida News (2010) 2pp.; http://news.ufl.edu/2010/03/31/antenna-pill-2/.

Intromedic, MicroCam Innovative Capsule Endoscope Pamphlet. (2006) 8 pp (http://www.intromedic.com/en/product/productinfo.asp) (8 pages).

ISFET—Ion Sensitive Field-Effect Transistor; Microsens S.A. pdf document. First cited by Examiner in Office Action dated Jun. 13 (2011) for U.S. Appl. No. 12/238,345; 4pp.

Jung, S. "Dissolvable 'Transient Electronics' Will Be Good for Your Body and the Environment" MedGadget; Oct. 1 (2012); Onlne website: http://medgadget.com/2012/10/dissolvable-transient-electronics-will-be-good-for-your-body-and-the-environment.html; downloaded Oct. 24, 2012; 4 pp.

Juvenile Diabetes Research Foundation International (JDRF), "Artificial Pancreas Project" (2010); http://www.artificialpancreasproject.com/; 3 pp.

Kamada K., "Electrophoretic deposition assisted by soluble anode" Materials Letters 57 (2003) 2348-2351.

Kendle, Earl R. and Morris, Larry A., "Preliminary Studies in the Development of a Gastric Battery for Fish" (1964). Nebraska Game and Parks Commission White Papers, Conference Presentations, & Manuscripts. Paper 22. p. 1-6.

Kim et al., "A Semi-Interpenetrating Network System for a Polymer Membrane"; Eur. Polym. J. vol. 33 No. 7; pp. 1009-1014 (1997). Li, P-Y, et al. "An electrochemical intraocular drug delivery device", Sensors and Actuators A 143 (2008) p. 41-48.

Lifescan, "OneTouch UltraLink<sup>TM</sup>" http://www.lifescan.com/products/meters/ultralink (2010) 2 pp.

Mackay et al., "Radio Telemetering from within the Body" Inside Information is Revealed by Tiny Transmitters that can be Swallowed or Implanted in Man or Animal Science (1991) 1196-1202; 134; American Association for the Advancement of Science, Washington D.C.

Mackay et al., "Endoradiosonde" Nature, (1957) 1239-1240, 179 Nature Publishing Group.

McKenzie et al., "Validation of a new telemetric core temperature monitor" J. Therm. Biol. (2004) 29(7-8):605-11.

Medtronic, "CareLink Therapy Management Software for Diabetes" (2010); https://carelink.minimed.com/patient/entry.jsp?bhcp=1; 1 pp.

Medfronic, "Carelink<sup>TM</sup> USB" (2008) http://www.medtronicdiabetes.com/pdf/carelink\_usb\_factsheet.pdf 2pp.

Medtronic "The New MiniMed Paradigm® REAL-Time Revel™ System" (2010) http://www.medtronicdiabetes.com/products/index. html; 2 pp.

Medtronic, "MINI MED Paradigm ® Revel <sup>TM</sup> Insulin Pump" (2010) http://www.medtronicdiabetes.com/products/insulinpumps/index.html; 2 pp.

Medtronic, Mini Med Paradigm TM VeoTM System: Factsheet (2010). http://www.medtronic-diabetes.com.au/downloads/Paradigm%20Veo% 20Factsheet.pdf; 4 pp.

Melanson, "Walkers swallow RFID pills for science" Engadget (2008); http://www.engadget.com/2008/07/29/walkers-swallow-rfid-pills-for-science/ (1 page).

Minimitter Co. Inc. "Actiheart" Traditional 510(k) Summary. Sep. 27, (2005) (8 pages).

Minimitter Co. Inc. Noninvasive technology to help your studies succeed. Mini Mitter.com Mar. 31, (2009) (4 pages).

Mini Mitter Co, Inc. 510(k) Premarket Notification Mini-Logger for Diagnostic Spirometer. 9-21 (1999) (9 pages).

Mini Mitter Co, Inc. 510(k) Premarket Notification for VitalSense. Apr. 22, 2004 (11 pages).

Minimitter Co. Inc. VitalSense Integrated Physiological Monitoring System. Product Description. (2005) (4 pages).

Minimitter Co. Inc. VitalSense Wireless Vital Signs Monitoring. Temperatures.com Mar. 31, 2009 (3 pages).

Mojaverian et al., "Estimation of gastric residence time of the Heidelberg capsule in humans: effect of varying food composition" Gastroenterology (1985) 89:(2): 392-7.

O'Brien et al., "The Production and Characterization of Chemically Reactive Porous Coatings of Zirconium Via Unbalanced Magnetron Sputtering" Surface and Coatings Technology (1996) 86-87; 200-

Park, "Medtronic to Buy MiniMed for \$3.7 Billion" (2001) HomeCare; http://homecaremag.com/mag/medical\_medtronic\_buy\_minimedl; 2 pp.

Philips Respironics Products, Noninvasive Technology to Help Your Studies Succeed. 510 (k) Permanent Notification for Vital Sense. Apr. 22, 2004; http/minimitter.com/products.cfm.

Radio Antennae, http://www.erikdeman.de/html/sail018h.htm; (2008) 5 pages.

"RFID "pill" monitors marchers" RFID News (2008) http://www.rfidnews.org/2008/07/23/rfid-pill-monitors-marchers/ (4 pages). Rolison et al., "Electrically conductive oxide aerogels: new materials in electrochemistry" J. Mater. Chem. (2001) 1, 963-980.

### (56) References Cited

### OTHER PUBLICATIONS

Roulstone, et al., "Studies on Polymer Latex Films: I. A study of latex film morphology" Polymer International 24 (1991) pp. 87-94. Sanduleanu et al., "Octave tunable, highly linear, RC-ring oscillator with differential fine-coarse tuning, quadrature outputs and amplitude control for fiber optic transceivers" (2002) IEEE MTT-S International Microwave Symposium Digest 545-8.

Santini, J.T. et al, "Microchips as controlled drug delivery-devices", Agnew. Chem. Int. Ed. (2000), vol. 39, p. 2396-2407.

"SensiVida minimally invasive clinical systems" Investor Presentation Oct. (2009) 28pp; http://www.sensividamedtech.com/SensiVidaGeneralOctober09.pdf; pp. 1-28.

Shawgo, R.S. et al. "BioMEMS from drug delivery", Current Opinion in Solid State and Material Science 6 (2002), p. 329-334. Shin et al., "A Simple Route to Metal Nanodots and Nanoporous Metal Films"; Nano Letters, vol. 2, No. 9 (2002) pp. 933-936.

Shrivas et al., "A New Platform for Bioelectronics-Electronic Pill", Cummins College, (2010).; http://www.cumminscollege.org/downloads/electronics\_and\_telecommunication/Newsletters/Current% 20Newsletters.pdf; First cited in third party client search conducted by Patent Eagle Search May 18, 2010 (2010); pp. 11-12.

"Smartlife awarded patent for knitted transducer" Innovation in Textiles News: http://www.innovationintextiles.com/articles/208. php; 2pp. (2009).

"The SmartPill Wireless Motility Capsule" SMARTPILL, The Measure of GI Health; (2010) http://www.smartpillcorp.com/index.cfm?pagepath=Products/The\_SmartPill\_Capsule&id=17814 (1 page). Solanas et al., "RFID Technology for the Health Care Sector" Recent Patents on Electrical Engineering (2008) 1, 22-31.

Soper, S.A. et al. "Bio-Mems Technologies and Applications", Chapter 12, "MEMS for Drug Delivery", p. 325-346 (2007).

Swedberg, "University Team Sees Ingestible RFID Tag as a Boon to Clinical Trials" RFID Journal (2010) Apr. 27; http://www.rfidjournal.com/article/view/7560/1 3pp.

Tajalli et al., "Improving the power-delay performance in subthreshold source-coupled logic circuits" Integrated Circuit and System Design. Power and Timing Modeling, Optimization and Simulation, Springer Berlin Heidelberg (2008) 21-30.

Target Innovations, Tablet Metal Detector, https://web. arch ive. org/web/20 130215063351 /http://www.metaldetectorindia.com/tablet -metal-detector. html, Feb. 15, 2013.

TargetPharmaceutical Metal Detector, Feb. 15, 2013 downloaded from Target Innovations, Tablet Metal Detector, Feb. 15, 2013.

Tatbul et al., "Confidence-based data management for personal area sensor networks" ACM International Conference Proceeding Series (2004) 72; 3 pages.

Tierney, M.J. et al "Electroreleasing Composite Membranes for Delivery of Insulin and other Biomacromolecules", J. Electrochem. Soc., vol. 137, No. 6, Jun. (1990), p. 2005-2006.

Trutag Technologies, Inc., Spectral Microtags for Authentication and Anti-Counterfeiting; "Product Authentication and Brand Protection Solutions"; http://www.trutags.com/; downloaded Feb. 12, 2013; 1 pp.

Walkey, "MOSFET Structure and Processing"; 97.398\* Physical Electronics Lecture 20; 24 pp.

Wang, X. et al "Resistance to Tracking and Erosion of Silicone Rubber Material under Various Types of Precipitation", Jpn. J. Appl. Phys. vol. 38 (1999) pp. 5170-5175.

Watson, et al., "Determination of the relationship between the pH and conductivity of gastric juice" Physiol Meas. 17 (1996) pp. 21-27.

Winter, J. et al. "The material properties of gelatin gels"; USA Ballistic Research Laboratories, Mar. (1975), p. 1-157.

Wongmanerod et al., "Determination of pore size distribution and surface area of thin porous silicon layers by spectroscopic ellipsometry" Applied Surface Science 172 (2001) 117-125.

Xiaoming et al., "A telemedicine system for wireless home healthcare based on bluetooth and the internet" Telemedicine Journal and e-health (2004) 10(S2): S110-6.

Yang et al., "Fast-switching frequency synthesizer with a discriminatoraided phase detector" IEEE Journal of Solid-State Circuits (2000) 35(10): 1445-52.

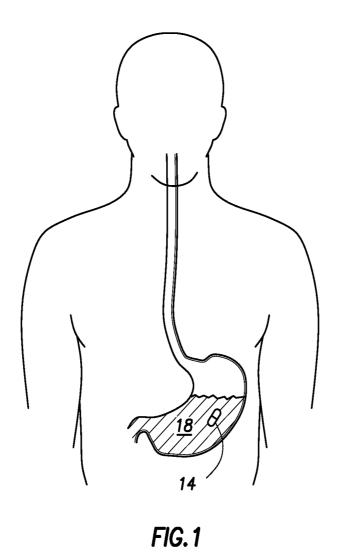
Yao et al., "Low Power Digital Communication in Implantable Devices Using vol. Conduction of Biological Tissues" Proceedings of the 28th IEEE, EMBS Annual International Conference, Aug. 30-Sep. 3, (2006); pp. 6249-6252.

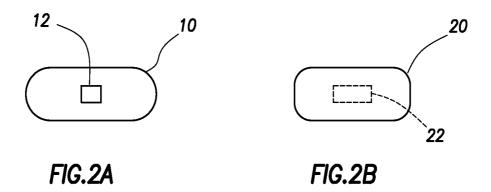
Youtube video Pharmaceutical Metal Detector/Tablet Metal Detector/ Capsule Metal Detector/ Dry Fruits; https://www.youtube.com/watch?v=I0126txam\_s, May 12, 2012.

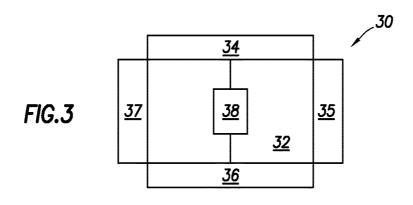
Zimmerman, "Personal Area Networks: Near-field intrabody communication" IBM Systems Journal (1996) 35 (3-4):609-17.

Zworkin, "A Radio Pill" Nature, (1957) 898, 179 Nature Publishing Group.

<sup>\*</sup> cited by examiner







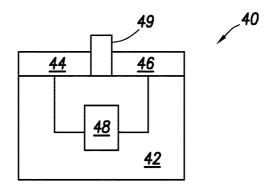


FIG.4

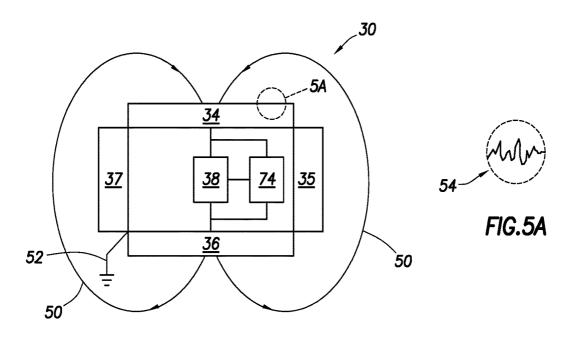
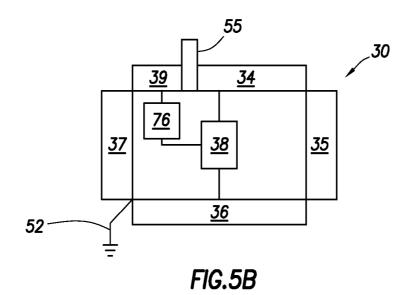
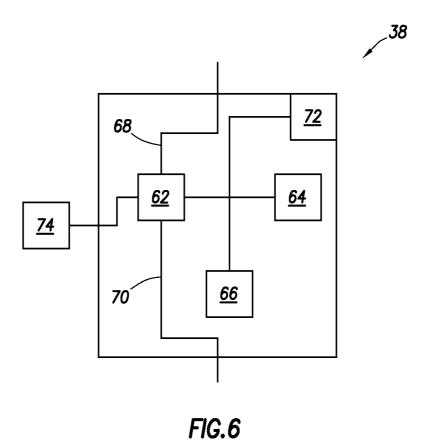


FIG.5





### **COMMUNICATION SYSTEM WITH** PARTIAL POWER SOURCE

### CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation application claiming priority under 35 U.S.C. § 120 to U.S. patent application Ser. No. 14/865,508, filed Sep. 25, 2015, now U.S. Pat. No. 9,649,066, which is a continuation application claiming  $^{10}$ priority under 35 U.S.C. § 120 to U.S. patent application Ser. No. 14/341,639, filed Jul. 25, 2014, now U.S. Patent Application Publication No. 2015/0080680 which is a continuation application claiming priority under 35 U.S.C. § 120 to U.S. patent application Ser. No. 13/153,312, filed Jun. 3, 2011, now U.S. Pat. No. 8,816,847, which is a continuation application claiming priority under 35 U.S.C. § 120 to U.S. patent application Ser. No. 12/564,017, filed Sep. 21, 2009, now U.S. Pat. No. 7,978,064, the entire disclosures of which are hereby herein incorporated by reference herein.

#### **FIELD**

The present invention is related to systems for detection of an event. More specifically, the present disclosure 25 includes a system that includes a partial power source that can be activated when in contact with conductive liquid and is capable of controlling conductance to mark an event.

### **BACKGROUND**

Ingestible devices that include electronic circuitry have been proposed for use in a variety of different medical applications, including both diagnostic and therapeutic applications. These devices typically require an internal 35 power supply for operation. Examples of such ingestible devices are ingestible electronic capsules which collect data as they pass through the body, and transmit the data to an external receiver system. An example of this type of electronic capsule is an in-vivo video camera. The swallowable 40 capsule includes a camera system and an optical system for imaging an area of interest onto the camera system. The transmitter transmits the video output of the camera system and the reception system receives the transmitted video device, which has an internal and self contained power source, which obtains images from within body lumens or cavities. The electronic circuit components of the device are enclosed by an inert indigestible housing (e.g. glass housing) that passes through the body internally. Other examples 50 include an ingestible data recorder capsule medical device. The electronic circuits of the disclosed device (e.g. sensor, recorder, battery etc.) are housed in a capsule made of inert materials.

In other examples, fragile radio frequency identification 55 (RFID) tags are used in drug ingestion monitoring applications. In order for the RFID tags to be operational, each requires an internal power supply. The RFID tags are antenna structures that are configured to transmit a radiofrequency signal through the body.

The problem these existing devices pose is that the power source is internal to device and such power sources are costly to produce and potentially harmful to the surrounding environment if the power source leaks or is damaged. Additionally, having antennas extending from the device is 65 a concern as related to the antennas getting damaged or causing a problem when the device is used in-vivo. There-

fore, what is needed is suitable system with circuitry that eliminates the need for an internal power source and anten-

### **SUMMARY**

The present disclosure includes a system for producing a unique signature that indicates the occurrence of an event. The system includes circuitry and components that can be placed within certain environments that include a conducting fluid. One example of such an environment is inside a container that houses the conducting fluid, such as a sealed bag with a solution, which includes an IV bag. Another example is within the body of a living organism, such as an animal or a human. The systems are ingestible and/or digestible or partially digestible. The system includes dissimilar materials positioned on the framework such that when a conducting fluid comes into contact with the dissimilar materials, a voltage potential difference is created. The voltage potential difference, and hence the voltage, is used to power up control logic that is positioned within the framework. Ions or current flows from the first dissimilar material to the second dissimilar material via the control logic and then through the conducting fluid to complete a circuit. The control logic controls the conductance between the two dissimilar materials and, hence, controls or modulates the conductance.

As the ingestible circuitry is made up of ingestible, and even digestible, components, the ingestible circuitry results in little, if any, unwanted side effects, even when employed in chronic situations. Examples of the range of components that may be included are: logic and/or memory elements; effectors; a signal transmission element; and a passive element, such as a resistor or inductor. The one or more components on the surface of the support may be laid out in any convenient configuration. Where two or more components are present on the surface of the solid support, interconnects may be provided. All of the components and the support of the ingestible circuitry are ingestible, and in certain instances digestible or partially digestible.

### BRIEF DESCRIPTION OF THE FIGURES

FIG. 1 shows a pharmaceutical product with an event output. Other examples include an ingestible imaging 45 indicator system according to the teaching of the present invention, wherein the product and the event indicator system combination are within the body.

> FIG. 2A shows the pharmaceutical product of FIG. 1 with the event indicator system on the exterior of the pharmaceutical product.

FIG. 2B shows the pharmaceutical product of FIG. 1 with the event indicator system positioned inside the pharmaceutical product.

FIG. 3 is a block diagram representation of one aspect of the event indicator system with dissimilar metals positioned on opposite ends.

FIG. 4 is a block diagram representation of another aspect of the event indicator system with dissimilar metals positioned on the same end and separated by a non-conducting 60 material.

FIG. 5 shows ionic transfer or the current path through a conducting fluid when the event indicator system of FIG. 3 is in contact with conducting liquid and in an active state.

FIG. 5A shows an exploded view of the surface of dissimilar materials of FIG. 5.

FIG. 5B shows the event indicator system of FIG. 5 with a pH sensor unit.

FIG. 6 is a block diagram illustration of one aspect of the control device used in the system of FIGS. 3 and 4.

### DETAILED DESCRIPTION

The present disclosure includes multiple embodiments for indicating the occurrence of an event. As described in more detail below, a system of the present invention is used with a conducting fluid to indicate the event marked by contact between the conducting fluid and the system. For example, the system of the present disclosure may be used with pharmaceutical product and the event that is indicated is when the product is taken or ingested. The term "ingested" or "ingest" or "ingesting" is understood to mean any introduction of the system internal to the body. For example, 15 ingesting includes simply placing the system in the mouth all the way to the descending colon. Thus, the term ingesting refers to any instant in time when the system is introduced to an environment that contains a conducting fluid. Another example would be a situation when a non-conducting fluid 20 is mixed with a conducting fluid. In such a situation the system would be present in the non-conduction fluid and when the two fluids are mixed, the system comes into contact with the conducting fluid and the system is activated. Yet another example would be the situation when the 25 presence of certain conducting fluids needed to be detected. In such instances, the presence of the system, which would be activated, within the conducting fluid could be detected and, hence, the presence of the respective fluid would be detected.

Referring again to the instance where the system is used with the product that is ingested by the living organism, when the product that includes the system is taken or ingested, the device comes into contact with the conducting liquid of the body. When the system of the present invention 35 comes into contact with the body fluid, a voltage potential is created and the system is activated. A portion of the power source is provided by the device, while another portion of the power source is provided by the conducting fluid, which is discussed in detail below.

Referring now to FIG. 1, an ingestible product 14 that includes a system of the present invention is shown inside the body. The product 14 is configured as an orally ingestible pharmaceutical formulation in the form of a pill or capsule. Upon ingestion, the pill moves to the stomach. Upon reaching the stomach, the product 14 is in contact with stomach fluid 18 and undergoes a chemical reaction with the various materials in the stomach fluid 18, such as hydrochloric acid and other digestive agents. The system of the present invention is discussed in reference to a pharmaceutical environment. However, the scope of the present invention is not limited thereby. The present invention can be used in any environment where a conducting fluid is present or becomes present through mixing of two or more components that result in a conducting liquid.

Referring now to FIG. 2A, a pharmaceutical product 10, similar to the product 14 of FIG. 1, is shown with a system 12, such as an ingestible event marker or an ionic emission module. The scope of the present invention is not limited by the shape or type of the product 10. For example, it will be 60 clear to one skilled in the art that the product 10 can be a capsule, a time-release oral dosage, a tablet, a gel cap, a sub-lingual tablet, or any oral dosage product that can be combined with the system 12. In the referenced embodiment, the product 10 has the system 12 secured to the 65 exterior using known methods of securing micro-devices to the exterior of pharmaceutical products. Example of meth-

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ods for securing the micro-device to the product is disclosed in U.S. Provisional Application No. 61/142,849 filed on Jan. 1, 2009 and entitled "HIGH-THROUGHPUT PRODUCTION OF INGESTIBLE EVENT MARKERS" as well as U.S. Provisional Application No. 61/177,611 filed on May 12, 2009 and entitled "INGESTIBLE EVENT MARKERS COMPRISING AN IDENTIFIER AND AN INGESTIBLE COMPONENT", the entire disclosure of each is incorporated herein by reference. Once ingested, the system 12 comes into contact with body liquids and the system 12 is activated. The system 12 uses the voltage potential difference to power up and thereafter modulates conductance to create a unique and identifiable current signature. Upon activation, the system 12 controls the conductance and, hence, current flow to produce the current signature.

There are various reasons for delaying the activation of the system 12. In order to delay the activation of the system 12, the system 12 may be coated with a shielding material or protective layer. The layer is dissolved over a period of time, thereby allowing the system 12 to be activated when the product 10 has reached a target location.

Referring now to FIG. 2B, a pharmaceutical product 20, similar to the product 14 of FIG. 1, is shown with a system 22, such as an ingestible event marker or an identifiable emission module. The scope of the present invention is not limited by the environment to which the system 22 is introduced. For example, the system 22 can be enclosed in a capsule that is taken in addition to/independently from the pharmaceutical product. The capsule may be simply a carrier for the system 22 and may not contain any product. Furthermore, the scope of the present invention is not limited by the shape or type of product 20. For example, it will be clear to one skilled in the art that the product 20 can be a capsule, a time-release oral dosage, a tablet, a gel capsule, a sublingual tablet, or any oral dosage product. In the referenced embodiment, the product 20 has the system 22 positioned inside or secured to the interior of the product 20. In one embodiment, the system 22 is secured to the interior wall of the product 20. When the system 22 is positioned inside a gel capsule, then the content of the gel capsule is a nonconducting gel-liquid. On the other hand, if the content of the gel capsule is a conducting gel-liquid, then in an alternative embodiment, the system 22 is coated with a protective cover to prevent unwanted activation by the gel capsule content. If the content of the capsule is a dry powder or microspheres, then the system 22 is positioned or placed within the capsule. If the product 20 is a tablet or hard pill, then the system 22 is held in place inside the tablet. Once ingested, the product 20 containing the system 22 is dissolved. The system 22 comes into contact with body liquids and the system 22 is activated. Depending on the product 20, the system 22 may be positioned in either a near-central or near-perimeter position depending on the desired activation delay between the time of initial ingestion and activation of 55 the system 22. For example, a central position for the system 22 means that it will take longer for the system 22 to be in contact with the conducting liquid and, hence, it will take longer for the system 22 to be activated. Therefore, it will take longer for the occurrence of the event to be detected.

Referring now to FIG. 3, in one embodiment, the systems 12 and 22 of FIGS. 2A and 2B, respectively, are shown in more detail as system 30. The system 30 can be used in association with any pharmaceutical product, as mentioned above, to determine when a patient takes the pharmaceutical product. As indicated above, the scope of the present invention is not limited by the environment and the product that is used with the system 30. For example, the system 30 may

be placed within a capsule and the capsule is placed within the conducting liquid. The capsule would then dissolve over a period of time and release the system 30 into the conducting liquid. Thus, in one embodiment, the capsule would contain the system 30 and no product. Such a capsule may then be used in any environment where a conducting liquid is present and with any product. For example, the capsule may be dropped into a container filled with jet fuel, salt water, tomato sauce, motor oil, or any similar product. Additionally, the capsule containing the system 30 may be ingested at the same time that any pharmaceutical product is ingested in order to record the occurrence of the event, such as when the product was taken.

In the specific example of the system 30 combined with the pharmaceutical product, as the product or pill is ingested, the system 30 is activated. The system 30 controls conductance to produce a unique current signature that is detected, thereby signifying that the pharmaceutical product has been taken. The system 30 includes a framework 32. The frame- 20 work 32 is a chassis for the system 30 and multiple components are attached to, deposited upon, or secured to the framework 32. In this embodiment of the system 30, a digestible material 34 is physically associated with the framework 32. The material 34 may be chemically deposited 25 on, evaporated onto, secured to, or built-up on the framework all of which may be referred to herein as "deposit" with respect to the framework 32. The material 34 is deposited on one side of the framework 32. The materials of interest that can be used as material 34 include, but are not limited to: Cu or CuI. The material 34 is deposited by physical vapor deposition, electrodeposition, or plasma deposition, among other protocols. The material 34 may be from about 0.05 to about 500 µm thick, such as from about 5 to about 100 μm thick. The shape is controlled by shadow mask deposition, or photolithography and etching. Additionally, even though only one region is shown for depositing the material, each system 30 may contain two or more electrically unique regions where the material 34 may be deposited, as desired.

At a different side, which is the opposite side as shown in FIG. 3, another digestible material 36 is deposited, such that materials 34 and 36 are dissimilar. Although not shown, the different side selected may be the side next to the side 45 selected for the material 34. The scope of the present invention is not limited by the side selected and the term "different side" can mean any of the multiple sides that are different from the first selected side. Furthermore, even though the shape of the system is shown as a square, the 50 shape maybe any geometrically suitable shape. Material 34 and 36 are selected such that they produce a voltage potential difference when the system 30 is in contact with conducting liquid, such as body fluids. The materials of interest for material 36 include, but are not limited to: Mg, Zn, or 55 other electronegative metals. As indicated above with respect to the material 34, the material 36 may be chemically deposited on, evaporated onto, secured to, or built-up on the framework. Also, an adhesion layer may be necessary to help the material 36 (as well as material 34 when needed) to 60 adhere to the framework 32. Typical adhesion layers for the material 36 are Ti, TiW, Cr or similar material. Anode material and the adhesion layer may be deposited by physical vapor deposition, electrodeposition or plasma deposition. The material 36 may be from about 0.05 to about 500 65 μm thick, such as from about 5 to about 100 μm thick. However, the scope of the present invention is not limited by

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the thickness of any of the materials nor by the type of process used to deposit or secure the materials to the framework 32.

According to the disclosure set forth, the materials 34 and 36 can be any pair of materials with different electrochemical potentials. Additionally, in the embodiments wherein the system 30 is used in-vivo, the materials 34 and 36 may be vitamins that can be absorbed. More specifically, the materials 34 and 36 can be made of any two materials appropriate for the environment in which the system 30 will be operating. For example, when used with an ingestible product, the materials 34 and 36 are any pair of materials with different electrochemical potentials that are ingestible. An illustrative example includes the instance when the system 30 is in contact with an ionic solution, such as stomach acids. Suitable materials are not restricted to metals, and in certain embodiments the paired materials are chosen from metals and non-metals, e.g., a pair made up of a metal (such as Mg) and a salt (such as CuCl or CuI). With respect to the active electrode materials, any pairing of substances-metals, salts, or intercalation compounds—with suitably different electrochemical potentials (voltage) and low interfacial resistance are suitable.

Materials and pairings of interest include, but are not limited to, those reported in Table 1 below. In one embodiment, one or both of the metals may be doped with a non-metal, e.g., to enhance the voltage potential created between the materials as they come into contact with a conducting liquid. Non-metals that may be used as doping agents in certain embodiments include, but are not limited to: sulfur, iodine and the like. In another embodiment, the materials are copper iodine (CuI) as the anode and magnesium (Mg) as the cathode. Embodiments of the present invention use electrode materials that are not harmful to the human body.

TABLE 1

	Anode	Cathode
Metals	Magnesium, Zinc Sodium (†), Lithium (†) Iron	
Salts	N/	Copper salts: iodide, chloride, bromide, sulfate, formate, (other anions possible) Fe <sup>3+</sup> salts: e.g. orthophosphate, pyrophosphate, (other anions possible) Oxygen (††) on platinum, gold or other catalytic surfaces
Intercalation compounds	Graphite with Li, K, Ca, Na, Mg	Vanadium oxide Manganese oxide

Thus, when the system 30 is in contact with the conducting liquid, a current path, an example is shown in FIG. 5, is formed through the conducting liquid between material 34 and 36. A control device 38 is secured to the framework 32 and electrically coupled to the materials 34 and 36. The control device 38 includes electronic circuitry, for example control logic that is capable of controlling and altering the conductance between the materials 34 and 36.

The voltage potential created between the materials 34 and 36 provides the power for operating the system as well as produces the current flow through the conducting fluid and the system. In one embodiment, the system operates in direct current mode. In an alternative embodiment, the system controls the direction of the current so that the direction of current is reversed in a cyclic manner, similar to alternating current. As the system reaches the conducting fluid or the electrolyte, where the fluid or electrolyte com-

ponent is provided by a physiological fluid, e.g., stomach acid, the path for current flow between the materials **34** and **36** is completed external to the system **30**; the current path through the system **30** is controlled by the control device **38**. Completion of the current path allows for the current to flow 5 and in turn a receiver, not shown, can detect the presence of the current and recognize that the system **30** has been activate and the desired event is occurring or has occurred.

In one embodiment, the two materials 34 and 36 are similar in function to the two electrodes needed for a direct 10 current power source, such as a battery. The conducting liquid acts as the electrolyte needed to complete the power source. The completed power source described is defined by the physical chemical reaction between the materials 34 and 36 of the system 30 and the surrounding fluids of the body. 15 The completed power source may be viewed as a power source that exploits reverse electrolysis in an ionic or a conduction solution such as gastric fluid, blood, or other bodily fluids and some tissues. Additionally, the environment may be something other than a body and the liquid may 20 be any conducting liquid. For example, the conducting fluid may be salt water or a metallic based paint.

In certain embodiments, these two materials are shielded from the surrounding environment by an additional layer of material. Accordingly, when the shield is dissolved and the 25 two dissimilar materials are exposed to the target site, a voltage potential is generated.

In certain embodiments, the complete power source or supply is one that is made up of active electrode materials, electrolytes, and inactive materials, such as current collectors, packaging, etc. The active materials are any pair of materials with different electrochemical potentials. Suitable materials are not restricted to metals, and in certain embodiments the paired materials are chosen from metals and non-metals, e.g., a pair made up of a metal (such as Mg) and 35 a salt (such as CuI). With respect to the active electrode materials, any pairing of substances—metals, salts, or intercalation compounds—with suitably different electrochemical potentials (voltage) and low interfacial resistance are suitable.

A variety of different materials may be employed as the materials that form the electrodes. In certain embodiments, electrode materials are chosen to provide for a voltage upon contact with the target physiological site, e.g., the stomach, sufficient to drive the system of the identifier. In certain 45 embodiments, the voltage provided by the electrode materials upon contact of the metals of the power source with the target physiological site is 0.001 V or higher, including 0.01 V or higher, such as 0.1 V or higher, e.g., 0.3 V or higher, including 0.5 volts or higher, and including 1.0 volts or 50 higher, where in certain embodiments, the voltage ranges from about 0.001 to about 10 volts, such as from about 0.01 to about 10 V.

Referring again to FIG. 3, the materials 34 and 36 provide the voltage potential to activate the control device 38. Once 55 the control device 38 is activated or powered up, the control device 38 can alter conductance between the materials 34 and 36 in a unique manner. By altering the conductance between materials 34 and 36, the control device 38 is capable of controlling the magnitude of the current through 60 the conducting liquid that surrounds the system 30. This produces a unique current signature that can be detected and measured by a receiver (not shown), which can be positioned internal or external to the body. In addition to controlling the magnitude of the current path between the 65 materials, non-conducting materials, membrane, or "skirt" are used to increase the "length" of the current path and,

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hence, act to boost the conductance path, as disclosed in the U.S. patent application Ser. No. 12/238,345 entitled, "In-Body Device with Virtual Dipole Signal Amplification" filed Sep. 25, 2008, the entire content of which is incorporated herein by reference. Alternatively, throughout the disclosure herein, the terms "non-conducting material", "membrane", and "skirt" are interchangeably with the term "current path extender" without impacting the scope or the present embodiments and the claims herein. The skirt, shown in portion at 35 and 37, respectively, may be associated with, e.g., secured to, the framework 32. Various shapes and configurations for the skirt are contemplated as within the scope of the present invention. For example, the system 30 may be surrounded entirely or partially by the skirt and the skirt maybe positioned along a central axis of the system 30 or off-center relative to a central axis. Thus, the scope of the present invention as claimed herein is not limited by the shape or size of the skirt. Furthermore, in other embodiments, the materials 34 and 36 may be separated by one skirt that is positioned in any defined region between the materials 34 and 36.

Referring now to FIG. 4, in another embodiment, the systems 12 and 22 of FIGS. 2A and 2B, respectively, are shown in more detail as system 40. The system 40 includes a framework 42. The framework 42 is similar to the framework 32 of FIG. 3. In this embodiment of the system 40, a digestible or dissolvable material 44 is deposited on a portion of one side of the framework 42. At a different portion of the same side of the framework 42, another digestible material 46 is deposited, such that materials 44 and 46 are dissimilar. More specifically, material 44 and 46 are selected such that they form a voltage potential difference when in contact with a conducting liquid, such as body fluids. Thus, when the system 40 is in contact with and/or partially in contact with the conducting liquid, then a current path, an example is shown in FIG. 5, is formed through the conducting liquid between material 44 and 46. A control device 48 is secured to the framework 42 and electrically coupled to the materials 44 and 46. The control device 48 includes electronic circuitry that is capable of controlling part of the conductance path between the materials 44 and 46. The materials 44 and 46 are separated by a nonconducting skirt 49. Various examples of the skirt 49 are disclosed in U.S. Provisional Application No. 61/173,511 filed on Apr. 28, 2009 and entitled "HIGHLY RELIABLE INGESTIBLE EVENT MARKERS AND METHODS OF USING SAME" and U.S. Provisional Application No. 61/173,564 filed on Apr. 28, 2009 and entitled "INGEST-IBLE EVENT MARKERS HAVING SIGNAL AMPLIFI-ERS THAT COMPRISE AN ACTIVE AGENT"; as well as U.S. application Ser. No. 12/238,345 filed Sep. 25, 2008 and entitled "IN-BODY DEVICE WITH VIRTUAL DIPOLE SIGNAL AMPLIFICATION"; the entire disclosure of each is incorporated herein by reference.

Once the control device 48 is activated or powered up, the control device 48 can alter conductance between the materials 44 and 46. Thus, the control device 48 is capable of controlling the magnitude of the current through the conducting liquid that surrounds the system 40. As indicated above with respect to system 30, a unique current signature that is associated with the system 40 can be detected by a receiver (not shown) to mark the activation of the system 40. In order to increase the "length" of the current path the size of the skirt 49 is altered. The longer the current path, the easier it may be for the receiver to detect the current.

Referring now to FIG. 5, the system 30 of FIG. 3 is shown in an activated state and in contact with conducting liquid.

The system 30 is grounded through ground contact 52. The system 30 also includes a sensor module 74, which is described in greater detail with respect to FIG. 6. Ion or current paths 50 between material 34 to material 36 and through the conducting fluid in contact with the system 30. 5 The voltage potential created between the material 34 and 36 is created through chemical reactions between materials 34/36 and the conducting fluid. FIG. 5A shows an exploded view of the surface of the material 34. The surface of the material 34 is not planar, but rather an irregular surface. The 10 irregular surface increases the surface area of the material and, hence, the area that comes in contact with the conducting fluid.

In one embodiment, at the surface of the material 34, there is chemical reaction between the material 34 and the sur- 15 rounding conducting fluid such that mass is released into the conducting fluid. The term "mass" as used herein refers to protons and neutrons that form a substance. One example includes the instant where the material is CuCl and when in contact with the conducting fluid, CuCl becomes Cu (solid) 20 and Cl<sup>-</sup> in solution. The flow of ions into the conduction fluid is depicted by the ion paths 50. In a similar manner, there is a chemical reaction between the material 36 and the surrounding conducting fluid and ions are captured by the material 36. The release of ions at the material 34 and 25 capture of ion by the material 36 is collectively referred to as the ionic exchange. The rate of ionic exchange and, hence the ionic emission rate or flow, is controlled by the control device 38. The control device 38 can increase or decrease the rate of ion flow by altering the conductance, which alters the 30 impedance, between the materials 34 and 36. Through controlling the ion exchange, the system 30 can encode information in the ionic exchange process. Thus, the system 30 uses ionic emission to encode information in the ionic exchange.

The control device 38 can vary the duration of a fixed ionic exchange rate or current flow magnitude while keeping the rate or magnitude near constant, similar to when the frequency is modulated and the amplitude is constant. Also, the control device 38 can vary the level of the ionic 40 exchange rate or the magnitude of the current flow while keeping the duration near constant. Thus, using various combinations of changes in duration and altering the rate or magnitude, the control device 38 encodes information in the current flow or the ionic exchange. For example, the control 45 device 38 may use, but is not limited to any of the following techniques namely, Binary Phase-Shift Keying (PSK), Frequency modulation, Amplitude modulation, on-off keying, and PSK with on-off keying.

As indicated above, the various embodiments disclosed 50 herein, such as systems 30 and 40 of FIGS. 3 and 4, respectively, include electronic components as part of the control device 38 or the control device 48. Components that may be present include but are not limited to: logic and/or memory elements, an integrated circuit, an inductor, a 55 resistor, and sensors for measuring various parameters. Each component may be secured to the framework and/or to another component. The components on the surface of the support may be laid out in any convenient configuration. Where two or more components are present on the surface 60 of the solid support, interconnects may be provided.

As indicated above, the system, such as system 30 and 40, control the conductance between the dissimilar materials and, hence, the rate of ionic exchange or the current flow. Through altering the conductance in a specific manner the 65 system is capable of encoding information in the ionic exchange and the current signature. The ionic exchange or

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the current signature is used to uniquely identify the specific system. Additionally, the systems 30 and 40 are capable of producing various different unique exchanges or signatures and, thus, provide additional information. For example, a second current signature based on a second conductance alteration pattern may be used to provide additional information, which information may be related to the physical environment. To further illustrate, a first current signature may be a very low current state that maintains an oscillator on the chip and a second current signature may be a current state at least a factor of ten higher than the current state associated with the first current signature.

Referring now to FIG. 6, a block diagram representation of the control device 38 is shown. The device 30 includes a control module 62, a counter or clock 64, and a memory 66. Additionally, the device 38 is shown to include a sensor module 72 as well as the sensor module 74, which was referenced in FIG. 5. The control module 62 has an input 68 electrically coupled to the material 34 and an output 70 electrically coupled to the material **36**. The control module 62, the clock 64, the memory 66, and the sensor modules 72/74 also have power inputs (some not shown). The power for each of these components is supplied by the voltage potential produced by the chemical reaction between materials 34 and 36 and the conducting fluid, when the system 30 is in contact with the conducting fluid. The control module 62 controls the conductance through logic that alters the overall impedance of the system 30. The control module 62 is electrically coupled to the clock 64. The clock 64 provides a clock cycle to the control module 62. Based upon the programmed characteristics of the control module 62, when a set number of clock cycles have passed, the control module **62** alters the conductance characteristics between materials 34 and 36. This cycle is repeated and thereby the control 35 device 38 produces a unique current signature characteristic. The control module 62 is also electrically coupled to the memory 66. Both the clock 64 and the memory 66 are powered by the voltage potential created between the materials 34 and 36.

The control module 62 is also electrically coupled to and in communication with the sensor modules 72 and 74. In the embodiment shown, the sensor module 72 is part of the control device 38 and the sensor module 74 is a separate component. In alternative embodiments, either one of the sensor modules 72 and 74 can be used without the other and the scope of the present invention is not limited by the structural or functional location of the sensor modules 72 or 74. Additionally, any component of the system 30 may be functionally or structurally moved, combined, or repositioned without limiting the scope of the present invention as claimed. Thus, it is possible to have one single structure, for example a processor, which is designed to perform the functions of all of the following modules: the control module 62, the clock 64, the memory 66, and the sensor module 72 or 74. On the other hand, it is also within the scope of the present invention to have each of these functional components located in independent structures that are linked electrically and able to communicate.

Referring again to FIG. 6, the sensor modules 72 or 74 can include any of the following sensors: temperature, pressure, pH level, and conductivity. In one embodiment, the sensor modules 72 or 74 gather information from the environment and communicate the analog information to the control module 62. The control module then converts the analog information to digital information and the digital information is encoded in the current flow or the rate of the transfer of mass that produces the ionic flow. In another embodiment,

the sensor modules 72 or 74 gather information from the environment and convert the analog information to digital information and then communicate the digital information to control module 62. In the embodiment shown in FIG. 5, the sensor modules 74 is shown as being electrically coupled to 5 the material 34 and 36 as well as the control device 38. In another embodiment, as shown in FIG. 6, the sensor module 74 is electrically coupled to the control device 38 at connection 78. The connection 78 acts as both a source for power supply to the sensor module 74 and a communication 10 channel between the sensor module 74 and the control device 38

Referring now to FIG. 5B, the system 30 includes a pH sensor module 76 connected to a material 39, which is selected in accordance with the specific type of sensing function being performed. The pH sensor module 76 is also connected to the control device 38. The material 39 is electrically isolated from the material 34 by a non-conductive barrier 55. In one embodiment, the material 39 is platinum. In operation, the pH sensor module 76 uses the 20 voltage potential difference between the materials 34/36. The pH sensor module 76 measures the voltage potential difference between the material 34 and the material 39 and records that value for later comparison. The pH sensor between the material 39 and the material 36 and records that value for later comparison. The pH sensor module 76 calculates the pH level of the surrounding environment using the voltage potential values. The pH sensor module 76 provides that information to the control device 38. The 30 control device 38 varies the rate of the transfer of mass that produces the ionic transfer and the current flow to encode the information relevant to the pH level in the ionic transfer, which can be detected by a receiver (not shown). Thus, the system 30 can determine and provide the information related 35 to the pH level to a source external to the environment.

As indicated above, the control device 38 can be programmed in advance to output a pre-defined current signature. In another embodiment, the system can include a receiver system that can receive programming information 40 when the system is activated. In another embodiment, not shown, the switch 64 and the memory 66 can be combined into one device.

In addition to the above components, the system 30 may also include one or other electronic components. Electrical 45 components of interest include, but are not limited to: additional logic and/or memory elements, e.g., in the form of an integrated circuit; a power regulation device, e.g., battery, fuel cell or capacitor; a sensor, a stimulator, etc.; a signal transmission element, e.g., in the form of an antenna, 50 electrode, coil, etc.; a passive element, e.g., an inductor, resistor, etc.

In certain embodiments, the ingestible circuitry includes a coating layer. The purpose of this coating layer can vary, e.g., to protect the circuitry, the chip and/or the battery, or 55 any components during processing, during storage, or even during ingestion. In such instances, a coating on top of the circuitry may be included. Also of interest are coatings that are designed to protect the ingestible circuitry during storage, but dissolve immediately during use. For example, 60 coatings that dissolve upon contact with an aqueous fluid, e.g. stomach fluid, or the conducting fluid as referenced above. Also of interest are protective processing coatings that are employed to allow the use of processing steps that would otherwise damage certain components of the device. 65 For example, in embodiments where a chip with dissimilar material deposited on the top and bottom is produced, the

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product needs to be diced. However, the dicing process can scratch off the dissimilar material, and also there might be liquid involved which would cause the dissimilar materials to discharge or dissolve. In such instances, a protective coating on the materials prevents mechanical or liquid contact with the component during processing can be employed. Another purpose of the dissolvable coatings may be to delay activation of the device. For example, the coating that sits on the dissimilar material and takes a certain period of time, e.g., five minutes, to dissolve upon contact with stomach fluid may be employed. The coating can also be an environmentally sensitive coating, e.g., a temperature or pH sensitive coating, or other chemically sensitive coating that provides for dissolution in a controlled fashion and allows one to activate the device when desired. Coatings that survive the stomach but dissolve in the intestine are also of interest, e.g., where one desires to delay activation until the device leaves the stomach. An example of such a coating is a polymer that is insoluble at low pH, but becomes soluble at a higher pH. Also of interest are pharmaceutical formulation protective coatings, e.g., a gel cap liquid protective coating that prevents the circuit from being activated by liquid of the gel cap.

Identifiers of interest include two dissimilar electrochemimodule 76 also measures the voltage potential difference 25 cal materials, which act similar to the electrodes (e.g., anode and cathode) of a power source. The reference to an electrode or anode or cathode are used here merely as illustrative examples. The scope of the present invention is not limited by the label used and includes the embodiment wherein the voltage potential is created between two dissimilar materials. Thus, when reference is made to an electrode, anode, or cathode it is intended as a reference to a voltage potential created between two dissimilar materials.

> When the materials are exposed and come into contact with the body fluid, such as stomach acid or other types of fluid (either alone or in combination with a dried conductive medium precursor), a potential difference, that is, a voltage, is generated between the electrodes as a result of the respective oxidation and reduction reactions incurred to the two electrode materials. A voltaic cell, or battery, can thereby be produced. Accordingly, in embodiments of the invention, such power supplies are configured such that when the two dissimilar materials are exposed to the target site, e.g., the stomach, the digestive tract, etc., a voltage is generated.

> In certain embodiments, one or both of the metals may be doped with a non-metal, e.g., to enhance the voltage output of the battery. Non-metals that may be used as doping agents in certain embodiments include, but are not limited to: sulfur, iodine and the like.

> It is to be understood that this invention is not limited to particular embodiments or aspects described, as such may vary. It is also to be understood that the terminology used herein is for the purpose of describing particular embodiments only, and is not intended to be limiting, since the scope of the present invention will be limited only by the appended claims.

> Where a range of values is provided, it is understood that each intervening value, to the tenth of the unit of the lower limit unless the context clearly dictates otherwise, between the upper and lower limit of that range and any other stated or intervening value in that stated range, is encompassed within the invention. The upper and lower limits of these smaller ranges may independently be included in the smaller ranges and are also encompassed within the invention, subject to any specifically excluded limit in the stated range. Where the stated range includes one or both of the limits,

ranges excluding either or both of those included limits are also included in the invention.

Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this 5 invention belongs. Although any methods and materials similar or equivalent to those described herein can also be used in the practice or testing of the present invention, representative illustrative methods and materials are now described.

All publications and patents cited in this specification are herein incorporated by reference as if each individual publication or patent were specifically and individually indicated to be incorporated by reference and are incorporated herein by reference to disclose and describe the methods 15 and/or materials in connection with which the publications are cited. The citation of any publication is for its disclosure prior to the filing date and should not be construed as an admission that the present invention is not entitled to antedate such publication by virtue of prior invention. Further, 20 the dates of publication provided may be different from the actual publication dates which may need to be independently confirmed

It is noted that, as used herein and in the appended claims, the singular forms "a", "an", and "the" include plural 25 referents unless the context clearly dictates otherwise. It is further noted that the claims may be drafted to exclude any optional element. As such, this statement is intended to serve as antecedent basis for use of such exclusive terminology as "solely," "only" and the like in connection with the recitation of claim elements, or use of a "negative" limitation.

As will be apparent to those of skill in the art upon reading this disclosure, each of the individual embodiments described and illustrated herein has discrete components and features which may be readily separated from or combined 35 with the features of any of the other several embodiments without departing from the scope or spirit of the present invention. Any recited method can be carried out in the order of events recited or in any other order which is logically possible

Although the foregoing invention has been described in some detail by way of illustration and example for purposes of clarity of understanding, it is readily apparent to those of ordinary skill in the art in light of the teachings of this invention that certain changes and modifications may be 45 made thereto without departing from the spirit or scope of the appended claims.

Accordingly, the preceding merely illustrates the principles of the invention. It will be appreciated that those skilled in the art will be able to devise various arrangements 50 which, although not explicitly described or shown herein, embody the principles of the invention and are included within its spirit and scope. Furthermore, all examples and conditional language recited herein are principally intended to aid the reader in understanding the principles of the 55 invention and the concepts contributed by the inventors to furthering the art, and are to be construed as being without limitation to such specifically recited examples and conditions. Moreover, all statements herein reciting principles, aspects, and embodiments of the invention as well as spe- 60 cific examples thereof, are intended to encompass both structural and functional equivalents thereof. Additionally, it is intended that such equivalents include both currently known equivalents and equivalents developed in the future, i.e., any elements developed that perform the same function, 65 regardless of structure. The scope of the present invention, therefore, is not intended to be limited to the exemplary

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embodiments shown and described herein. Rather, the scope and spirit of present invention is embodied by the appended claims.

What is claimed is:

- 1. A composition, comprising:
- a pharmaceutical product comprising a pharmaceutically acceptable carrier; and
- an ingestible event marker coupled to the pharmaceutical product, wherein the ingestible event marker comprises:
  - a control device; and
  - a partial power source comprising first and second electrodes formed of dissimilar electrochemical materials configured to contact a conductive fluid and generate a voltage to energize the ingestible event marker:
  - wherein the control device of the energized ingestible event marker is configured to modulate current flow through the conductive fluid, the modulated current flow defining a unique current signature associated with the ingestible event marker;
- wherein the ingestible event marker of the composition is: coated with an external layer configured to delay activation of the ingestible event marker for a first period of time; and
  - positioned inside the pharmaceutically acceptable carrier, relative to a perimeter defined by the pharmaceutically acceptable carrier, to delay activation of the ingestible event marker for a second period of time, wherein the ingestible event marker is positioned proximate to the perimeter or proximate to a center of the pharmaceutically acceptable carrier based on an identity of the pharmaceutical product.
- 2. The composition according to claim 1, wherein the external layer comprises a homogeneous layer of a single material.
- 3. The composition according to claim 1, wherein the external layer comprises two or more distinct materials.
- **4**. The composition according to claim **3**, wherein the two or more distinct materials are present as a multilayer structure
- 5. The composition according to claim 1, wherein the external layer is configured to provide conductive fluid passage through the external layer after contact of the composition with the conductive fluid.
- **6**. The composition according to claim **1**, wherein the external layer is further configured to protect the ingestible event marker in a dicing process.
- 7. The composition according to claim 1, wherein the external layer is configured to dissolve in five or more minutes after being in contact with the conducting fluid.
- **8**. The composition according to claim **1**, wherein the external layer is environmentally sensitive.
- 9. The composition according to claim 8, wherein the external layer is temperature sensitive.
- 10. The composition according to claim 8, wherein the external layer is pH sensitive.
- 11. The composition according to claim 10, wherein the external layer is insoluble at a first pH and soluble at a second pH.
- 12. The composition according to claim 1, wherein the ingestible event marker is secured inside the pharmaceutically acceptable carrier.
- 13. The composition according to claim 1, wherein the first and second electrodes of the ingestible event marker are coated with the external layer.

- 14. The composition according to claim 1, wherein the pharmaceutical product further comprises a pharmaceutical material, and wherein the external layer is further configured to prevent the ingestible event marker from being activated by the pharmaceutical material.
- 15. The composition according to claim 1, wherein the ingestible event marker is positioned proximate to the perimeter or proximate to the center of the pharmaceutically acceptable carrier further based on a desired activation delay between a time of initial ingestion and activation of the ingestible event marker.
  - 16. A composition, comprising:
  - a pharmaceutical product comprising a pharmaceutically acceptable carrier; and
  - an ingestible event marker positioned inside the pharmaceutical product, wherein the ingestible event marker 15 comprises:
    - a partial power source comprising first and second electrodes formed of dissimilar electrochemical materials, wherein the partial power source is configured to generate a voltage to energize the ingestible event marker upon contact with a conductive fluid; and

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a control device configured to modulate current flow through the conductive fluid, wherein the modulated current flow defines a unique current signature;

wherein the ingestible event marker of the composition is:

coated with an external layer configured to at least one of delay activation of the ingestible event marker or protect the ingestible event marker; and

positioned relative to a perimeter defined by the pharmaceutically acceptable carrier to delay activation of the ingestible event marker, wherein the ingestible event marker is positioned proximate to the perimeter or proximate to a center of the pharmaceutically acceptable carrier based on an identity of the pharmaceutical product.

17. The composition according to claim 16, wherein the first and second electrodes of the ingestible event marker are coated with the external layer.

**18**. The composition according to claim **16**, wherein the external layer is environmentally sensitive.

\* \* \* \* \*



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### 摘要(译)

本发明的系统包括以不同材料形式的导电元件,电子部件和部分电源。 一旦与导电流体接触,就会产生电势并完成电源,从而激活系统。 电子 组件控制不同材料之间的电导,以产生独特的电流信号。 该系统可以用 于各种不同的应用中,包括作为可摄入标识符的组件,例如可以在可摄 入事件标记中找到的组件,例如,启用了药物信息学的药物组合物。

