

US009585593B2

(12) United States Patent

Fan et al.

(54) SIGNAL DISTRIBUTION FOR PATIENT-ELECTRODE MEASUREMENTS

- (76) Inventors: Chung Shing Fan, Toronto (CA); Joel Ironstone, Toronto (CA); Kenneth Carless Smith, Toronto (CA)
- (*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 1175 days.
- (21) Appl. No.: 13/508,655
- (22) PCT Filed: Nov. 18, 2010
- (86) PCT No.: PCT/AU2010/001552
 § 371 (c)(1),
 (2), (4) Date: Jan. 4, 2013
- (87) PCT Pub. No.: WO2011/060497PCT Pub. Date: May 26, 2011

(65) **Prior Publication Data**

US 2013/0102920 A1 Apr. 25, 2013

(30) Foreign Application Priority Data

Nov. 18, 2009 (AU) 2009905642

(51) Int. Cl.

A61B 5/05	(2006.01)
A61B 5/053	(2006.01)
	(Continued)

		(Commued)	
(52)	U.S. Cl.		

CPC *A61B 5/053* (2013.01); *A61B 5/0537* (2013.01); *A61B 5/4869* (2013.01); (Continued)

(10) Patent No.: US 9,585,593 B2

(45) **Date of Patent:** Mar. 7, 2017

(58) Field of Classification Search CPC A61B 5/053; A61B 5/0537; A61B 5/4869; A61B 5/4875; A61B 5/4878;

(Continued)

(56) **References Cited**

U.S. PATENT DOCUMENTS

3,316,896 A	5/1967	Thomasset
3,851,641 A	12/1974	Toole
5,051,011 11	(Con	tinued)

FOREIGN PATENT DOCUMENTS

CA	2231038	11/1999
CA	2638958	6/2000
	(Cor	tinued)

OTHER PUBLICATIONS

Abdullah M. Z.; Simulation of an inverse problem in electrical impedance tomography using resistance electrical network analogues; International Journal of Electrical Engineering Education; vol. 36, No. 4, pp. 311-324; Oct. 1999.

(Continued)

Primary Examiner — Devin Henson

(74) Attorney, Agent, or Firm — Knobbe Martens Olson & Bear LLP

(57) **ABSTRACT**

Apparatus for electrically connecting measurement apparatus to a biological subject, the apparatus including a signal delivery circuit including a current buffer having a current buffer input for receiving a signal from a signal source and a current buffer output for supplying a current to an electrode attached to the biological subject, and a voltage buffer having a voltage buffer input coupled to the current buffer output and a voltage buffer output for providing a voltage signal indicative of a voltage at the electrode, to a sensor.

24 Claims, 9 Drawing Sheets



(51) Int. Cl.

G01R 31/00	(2006.01)
G01R 27/02	(2006.01)
A61B 5/00	(2006.01)
H03F 3/50	(2006.01)
H03F 3/60	(2006.01)
G01R 1/30	(2006.01)

- (58) Field of Classification Search CPC ... A61B 5/7203; A61B 5/6843; A61B 5/6885; G01R 1/30; G01R 27/02; G01R 31/00; H03F 3/50; H03F 3/602; H03F 2200/261 See application file for complete search history.

(56) **References Cited**

U.S. PATENT DOCUMENTS

3.866.600	Α	2/1975	Rev
3,868,165	А	2/1975	Gonser
3.871.359	A	3/1975	Pacela
4.008.712	А	2/1977	Nyboer
4.034.854	A	7/1977	Bevilacqua
4.082.087	Ā	4/1978	Howson
4.121.575	Ā	10/1978	Mills et al.
4,144,878	Ā	3/1979	Wheeler
RE30.101	Ē	9/1979	Kubicek et al.
4 169 463	Ā	10/1979	Piquard
4 184 486	A	1/1980	Pana
4 233 987	Ā	11/1980	Feingold
4 291 708	A	9/1981	Frei et al
4 314 563	Ā	2/1982	Wheeler
4 353 372	Δ	10/1982	Aver
4 365 634	A	12/1982	Bare et al
4 407 288	Δ	10/1983	Langer et al
4 407 300	A	10/1983	Davis
4 450 527	7	5/108/	Sramek
4 4 58 604	Ā	7/108/	Sollish et al
4 486 835	Ā	12/1084	Boi et al
4,480,833	A	2/1025	Machida
4,537,203	A	0/1085	Fray et al
4,557,040	A	12/1085	Stoller et al
4,557,271	A	12/1985	Monoli
4,363,349	A	7/1086	Cook
4,002,338	A	10/1086	Drawn at al
4,017,939	A	1/1087	Diowii et al.
4,038,807	A	2/1087	Kydel Saala
4,040,734	A	3/1987	Civera et el
4,080,477	A	0/1907	Givens et al.
4,088,580	A	8/1987	Ko et al.
4,703,000	A	8/1988	Kroll et al.
4,793,302	A	5/1080	Iedner Van 11
4,852,008	A	5/1989	KIOII Carrana 1
4,836,214	A	6/1989	Sramek
4,890,630	A	1/1990	Kroll et al.
4,895,163	A	1/1990	Libke et al.
4,899,758	A	2/1990	Finkeistein et al.
4,905,705	A	3/1990	Kizakevich et al.
4,911,175	A	3/1990	Shizgal
4,922,911	A	5/1990	wada et al.
4,924,875	A	5/1990	Chamoun
4,942,880	A	//1990	Slovak
4,951,682	A	8/1990	Petre
4,981,141	A	1/1991	Segalowitz
5,020,541	A	6/1991	Marriott
5,025,784	A	6/1991	Shao et al.
5,063,937	A	11/1991	Ezenwa et al.
5,078,134	А	1/1992	Heilman et al.

5 006 501	2/1002	D 1
5,086,781 A	a 2/1992	Bookspan
5.101.828 A	4/1992	Welkowitz et al.
5 143 079 A	9/1992	Frei et al
5 184 624 4	2/1003	Brown et al
5,104,024 A	2/1993	
5,197,479 A	3/1993	Hubelbank et al.
5,199,432 A	4/1993	Quedens et al.
5.233.982 A	8/1993	Kohl
5 246 008 4	0/1003	Mueller
5,240,008 A	12/1002	Cianan at al
5,272,024 A	A 12/1993	Gisser et al.
5,280,429 A	a 1/1994	Withers
5.282.840 A	2/1994	Hudrlik
5 284 142 A	2/100/	Goble et al
5,205,102	4/1004	Dente et al.
3,303,192 A	4/1994	Bonte et al.
5,309,917 A	5/1994	Wang et al.
5,311,878 A	5/1994	Brown et al.
5 335 667 A	8/1994	Cha et al
5,353,007 1	10/1004	Changy at al
3,331,097 A	10/1994	Cheney et al.
5,353,802 A	A 10/1994	Ollmar
5,372,141 A	12/1994	Gallup et al.
5.381.333 A	1/1995	Isaacson et al.
5 300 110 4	2/1005	Cheney et al
5,550,110 A	£ (1005	Energy et al.
5,415,104 A	5/1995	Faupei
5,421,345 A	6/1995	Lekholm et al.
5,423,326 A	6/1995	Wang et al.
5 427 113 A	6/1995	Hiroshi et al
5 440 000 1	0/1005	Libko ot al
5,449,000 A	10/1005	
5,454,377 A	10/1995	Dzwonczyk et al.
5,465,730 A	11/1995	Zadehkoochak et al.
5.469.859 A	11/1995	Tsoglin et al.
5 503 157 A	4/1006	Sromek
5,505,157 A	4/1990	
5,505,209 A	4/1996	Reining
5,511,553 A	4/1996	Segalowitz
5.526.808 A	6/1996	Kaminsky
5 529 072 A	6/1996	Sramek
5 5 4 4 662	2/1006	Soulpier et el
5,544,002 A	0/1990	Sauliner et al.
5,557,242 A	a 9/1996	Wetherell
5,562,607 A	10/1996	Gyory
5.575.929 A	11/1996	Yu et al.
5 588 420 4	12/1006	Isaacson et al
5,500,429 A	2/1007	
5,011,351 A	A 3/1997	Sato et al.
5,615,689 A	4/1997	Kotler
5,626,146 A	5/1997	Barber et al.
5 704 355 A	1/1998	Bridges
5 730 136 1	3/1008	Laufor et al
5,750,150 A	a 3/1998	
5,732,710 A	A 3/1998	Rabinovich et al.
5,735,284 A	4/1998	Tsoglin et al.
5.746.214 A	5/1998	Brown et al.
5 750 150 A	6/1008	Masreliez
5700612 1	e/1009	Faldman
5,788,045 A	a 8/1998	Feldman
5,800,350 A	a 9/1998	Coppleson et al.
5,807,251 A	A 9/1998	Wang et al.
5.807.270 A	9/1998	Williams
5 807 272 A	0/1008	Kun et al
5,007,272 A	0/1008	Deselver
5,810,742 A	9/1998	Peariman
5,876,353 A	a 3/1999	Riff
5,919,142 A	. 7/1999	Boone et al.
5.947.910 A	9/1999	Zimmet
5 057 861 A	0/1000	Combs et al
5,957,801	10/1000	Combs et al.
5,964,703 A	10/1999	Goodman et al.
5,994,956 A	A 11/1999	Concorso
6,006,125 A	12/1999	Kelly et al.
6.011.992 A	1/2000	Hubbard et al.
6 015 389 A	1/2000	Brown
6 019 677 A	1/2000	Videina at al
0,018,077 A	A 1/2000	vidrine et al.
6,078,833 A	6/2000	T Tasa ala a m
6.101.413 A		Hueber
6.115.626 A	8/2000	Olson et al.
0,110,020 1	8/2000 9/2000	Olson et al. Whavne et al.
6 1 22 544 4	8/2000 9/2000	Olson et al. Whayne et al.
6,122,544 A	8/2000 9/2000 9/2000	Olson et al. Whayne et al. Organ
6,122,544 A 6,125,297 A	8/2000 9/2000 9/2000 9/2000 9/2000 9/2000	Olson et al. Whayne et al. Organ Siconolfi
6,122,544 A 6,125,297 A 6,129,666 A	x 8/2000 x 9/2000 x 9/2000 x 9/2000 x 9/2000 x 10/2000	Olson et al. Whayne et al. Organ Siconolfi DeLuca et al.
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A	x 8/2000 x 9/2000 x 9/2000 x 9/2000 x 10/2000 x 11/2000	Olson et al. Whayne et al. Organ Siconolfi DeLuca et al. Ubby
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A	A 8/2000 A 9/2000 A 9/2000 A 9/2000 A 10/2000 A 11/2000 A 11/2000	Olson et al. Whayne et al. Organ Siconolfi DeLuca et al. Ubby
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A 6,151,523 A	8/2000 8/2000 9/2000 9/2000 9/2000 9/2000 10/2000 11/2000 11/2000 11/2000	Olson et al. Whayne et al. Organ Siconolfi DeLuca et al. Ubby Ferrer et al.
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A 6,151,523 A 6,167,300 A	A 8/2000 A 9/2000 A 9/2000 A 9/2000 A 10/2000 A 11/2000 A 11/2000 A 12/2000	Olson et al. Olson et al. Organ Siconolfi DeLuca et al. Ubby Ferrer et al. Cherepenin et al.
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A 6,151,523 A 6,167,300 A 6,173,003 B	A 8/2000 A 9/2000 A 9/2000 A 10/2000 A 10/2000 A 11/2000 A 11/2000 A 12/2000 A 12/2000	Olson et al. Olson et al. Whayne et al. Organ Siconolfi DeLuca et al. Ubby Ferrer et al. Cherepenin et al. Whikehart et al.
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A 6,151,523 A 6,167,300 A 6,173,003 B 6,228 022 B	A 8/2000 A 9/2000 A 9/2000 A 9/2000 A 10/2000 A 11/2000 A 11/2000 A 12/2000 B1 1/2000 B1 1/2001	Olson et al. Olson et al. Whayne et al. Organ Siconolfi DeLuca et al. Ubby Ferrer et al. Cherepenin et al. Whikehart et al.
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A 6,151,523 A 6,167,300 A 6,173,003 B 6,228,022 B	x 8/2000 x 9/2000 x 9/2000 x 9/2000 x 10/2000 x 11/2000 x 12/2000 x 12/2000 x 12/2000 x 12/2000 x 12/2000	Olson et al. Olson et al. Organ Siconolfi DeLuca et al. Ubby Ferrer et al. Cherepenin et al. Whikehart et al. Friesem et al.
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A 6,151,523 A 6,167,300 A 6,173,003 B 6,228,022 B 6,228,033 B	x 8/2000 x 9/2000 x 9/2000 x 9/2000 x 9/2000 x 10/2000 x 11/2000 x 11/2000 x 12/2000 x 12/2000 x 12/2000 x 12/2000 x 12/2001 x 5/2001	Olson et al. Olson et al. Whayne et al. Organ Siconolfi DeLuca et al. Ubby Ferrer et al. Cherepenin et al. Whikehart et al. Friesem et al. Koobi
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A 6,151,523 A 6,167,300 A 6,173,003 B 6,228,022 B 6,228,033 B 6,233,473 B	x 8/2000 x 9/2000 x 9/2000 x 9/2000 x 9/2000 x 9/2000 x 10/2000 x 11/2000 x 12/2000 x 12/2001 x1 5/2001 x1 5/2001 x1 5/2001	Olson et al. Olson et al. Whayne et al. Organ Siconolfi DeLuca et al. Ubby Ferrer et al. Cherepenin et al. Whikehart et al. Friesem et al. Koobi Shepherd et al.
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A 6,151,523 A 6,167,300 A 6,173,003 B 6,228,022 B 6,228,033 B 6,228,033 B 6,233,473 B 6,26,886 B	x 8/2000 x 9/2000 x 9/2000 x 9/2000 x 10/2000 x 11/2000 x 11/2000 x 12/2000 x 12/2000 x 12/2001 x 5/2001 x 5/2001 x 5/2001	Olson et al. Olson et al. Organ Siconolfi DeLuca et al. Ubby Ferrer et al. Cherepenin et al. Whikehart et al. Friesem et al. Koobi Shepherd et al. Cherepenin et al.
6,122,544 A 6,125,297 A 6,129,666 A 6,142,949 A 6,151,523 A 6,173,003 B 6,228,022 B 6,228,033 B 6,233,473 B 6,236,886 B 6,236,886 B	x 8/2000 x 9/2000 x 9/2000 x 9/2000 x 9/2000 x 10/2000 x 11/2000 x 11/2000 x 12/2000 x 12/2000 x 12/2000 x 12/2001 x 5/2001 x 5/2001 x 5/2001	Olson et al. Olson et al. Whayne et al. Organ Siconolfi DeLuca et al. Ubby Ferrer et al. Cherepenin et al. Whikehart et al. Friesem et al. Koobi Shepherd et al. Cherepenin et al.

U.S. PATENT DOCUMENTS

6 253 100	D1	6/2001	Zhdanov
6 256 522	D1	7/2001	Cha
0,250,532	ы	7/2001	Cna
6,280,396	BI	8/2001	Clark
6,292,690	B1	9/2001	Petrucelli et al.
6,308,097	B1	10/2001	Pearlman
6.339.722	B1	1/2002	Heethaar et al.
6 354 006	BI	3/2002	Drinan et al
6,334,330		4/2002	Manian Ct al.
0,370,023	ы	4/2002	Mori
6,432,045	B2	8/2002	Lemperle et al.
6,459,930	B1	10/2002	Takehara et al.
6.469.732	B1	10/2002	Chang et al.
6 472 888	B2	10/2002	Oguma et al
6 406 725	D2 D2	12/2002	Vamada at al
6,490,723	DZ D1	12/2002	Ramada et al.
6,497,659	BI	12/2002	Kalert
6,501,984	B1	12/2002	Church et al.
6,511,438	B2	1/2003	Bernstein et al.
6.512.949	B1	1/2003	Combs et al.
6 516 218	B1	2/2003	Cheng et al
6 522 010	DI	2/2003	Gragami
6,522,910	DI	2/2003	
6,532,384	BI	3/2003	Fukuda
6,551,252	B2	4/2003	Sackner et al.
6,556,001	B1	4/2003	Wiegand et al.
6.560.480	B1	5/2003	Nachaliel et al.
6 561 986	B2	5/2003	Baura et al
6 564 070	D2 D1	5/2003	Comret al
0,304,079	DI	5/2003	
6,569,160	BI	5/2003	Goldin et al.
6,584,348	B2	6/2003	Glukhovsky
6,602,201	B1	8/2003	Hepp et al.
6.615.077	B1	9/2003	Zhu et al.
6 618 616	B2	9/2003	lijima et al
6 6 22 2 12	D2 D2	0/2003	Morray of al
6,025,512	D2 D2	9/2003	Meny et al.
6,625,487	B2	9/2003	Herleikson
6,631,292	B1	10/2003	Liedtk
6,633,777	B2	10/2003	Szopinski
6.636.754	B1	10/2003	Baura et al.
6 643 543	B2	11/2003	Takehara et al
6 658 206	D1	12/2003	Wong of al
6,038,290	DI	2/2003	
6,/14,813	B2	3/2004	Isnigooka et al.
6,714,814	B2	3/2004	Yamada et al.
6,723,049	B2	4/2004	Skladnev et al.
6.724.200	B2	4/2004	Fukuda
6 725 089	B2	4/2004	Komatsu et al
6 760 617	D2 D2	7/2004	Word of al
6,760,017	D2 D2	7/2004	Walu et al.
6,763,263	B2	7/2004	Gregory et al.
6,768,921	B2	7/2004	Organ et al.
6,788,966	B2	9/2004	Kenan et al.
6,790,178	B1	9/2004	Mault et al.
6.807.443	B2	10/2004	Keren
6 829 501	B2	12/2004	Nielsen et al
6 820 502	D2 D2	12/2004	A 14
0,829,303	D2	12/2004	
6,840,907	BI	1/2005	Brydon
6,845,264	B1	1/2005	Skladnev et al.
6,870,109	B1	3/2005	Villarreal
6.875.176	B2	4/2005	Mourad et al.
6 906 533	B1	6/2005	Yoshida
6 022 586	B7	7/2005	Davies
6,922,380	D2 D2	8/2005	W-ll-
6,936,012	BZ D2	8/2005	wens
6,940,286	B 2	9/2005	Wang et al.
RE38,879	E	11/2005	Goodman et al.
6,980,852	B2	12/2005	Jersey-Willuhn et al.
6.980.853	B2	12/2005	Miyoshi et al.
7 065 399	B2	6/2006	Nakada
7,005,555	D2 D2	7/2006	National and a
7,079,889	D2 D2	7/2000	INAKAUA
7,096,061	B 2	8/2006	Arad
7,122,012	B2	10/2006	Bouton et al.
7,130,680	B2	10/2006	Kodama et al.
7.132.611	B2	11/2006	Gregaard et al.
7 148 701	B2	12/2006	Park et al
7 140 572	B2	12/2006	Wong
7,177,373	D2 D2	1/2007	11 AUC 11 1 . 1
7,169,107	В2	1/2007	Jersey-Willuhn et al.
7,184,820	B2	2/2007	Jersey-Willuhn et al.
7,184.821	B2	2/2007	Belalcazar et al.
7 186 220	B2	3/2007	Stahmann et al
7 206 620	D2 D1	4/2007	Torlor
7,200,030	DI	4/2007	Tarter
7,212,852	B2	5/2007	Smith et al.

7,233,823	B2	6/2007	Simond et al.
7,251,524	B1	7/2007	Hepp et al.
7,288,943	B2	10/2007	Matthiessen et al.
7 313 435	э В2	12/2007 12/2007	Neverov et al. Nakada et al
7,317,161	B2 *	1/2008	Fukuda H01B 11/206
	DA	2/2000	174/36
7,336,992	B2 B2	2/2008	Shiokawa Woo et al
7,440,790	B2 B2	11/2008	Smith et al.
7,477,937	B2	1/2009	Iijima et al.
7,496,450	B2	2/2009	Ortiz Aleman et al.
7,499,745	B2	3/2009	Littrup et al.
D603,051 7,603,158	S B2	10/2009	Causevic et al.
7,603,171	B2 B2	10/2009	Eror et al.
7,628,761	B2	12/2009	Gozani et al.
7,638,341	B2	12/2009	Rubinsky et al.
7,657,292	B2	2/2010	Baker et al.
7,660,617	B2 D2	2/2010	Davis Min et al
7,700,872	B2 B2	5/2010	Garber et al
7,729,756	B2	6/2010	Mertelmeier et al.
7,733,224	B2	6/2010	Tran
7,749,013	B2	7/2010	Sato et al.
7,860,557	B2 B2	3/2010	Istvan et al.
D641 886	S	$\frac{3}{2011}$	Causevic et al
7,983,853	B2	7/2011	Wang et al.
D647,208	S	10/2011	Rothman et al.
8,055,335	B2	11/2011	Stylos
8,068,906	B2	11/2011	Chetham
8,172,702	B2 B2	7/2012	Koberison Johnson et al
8.233.974	B2	7/2012	Ward et al.
D669,186	S	10/2012	Gozani
D669,187	S	10/2012	Gozani
8,285,356	B2	10/2012	Bly et al.
D074,090 8467.865	ъ в2	6/2013	Gaw et al. Gregory et al
8,744.564	B2	6/2013	Ward et al.
D718,458	S	11/2014	Vosch et al.
D719,660	S	12/2014	Vosch et al.
D728,801	S	5/2015	Machon et al.
2001/0007030	A1 A1	7/2001	Kamada et al
2001/0020138	Al	9/2001	Ishigooka et al.
2001/0021799	A1	9/2001	Ohlsson et al.
2001/0025139	Al	9/2001	Pearlman
2001/0051774	AI	2/2002	Littrup et al. Walker et al
2002/0020138	Al	2/2002	Drinan et al.
2002/0022787	Al	2/2002	Takehara et al.
2002/0035334	Al	3/2002	Meij et al.
2002/0072686	Al	6/2002	Hoey et al.
2002/00/9910	A1 A1	7/2002	rukuda Kurihara
2002/0093992	Al	7/2002	Plangger
2002/0106681	A1	8/2002	Wexler et al.
2002/0109621	Al	8/2002	Khair et al.
2002/0111559	AI A1	8/2002	Kurata et al.
2002/0123034	Al	9/2002	Wexler et al.
2002/0161311	Al	10/2002	Ward et al.
2002/0193689	A1	12/2002	Bernstein et al.
2002/0194419	Al	12/2002	Rajput et al.
2003/0004403	AI A1	1/2003	Correct al
2003/0023184	Al	1/2003	Pitts-Crick et al.
2003/0028221	Al	2/2003	Zhu et al.
2003/0036713	A1	2/2003	Bouton et al.
2003/0050570	Al	3/2003	Kodama et al.
2003/0073916	Al	4/2003	Yonce
2003/0105410	AI A1	6/2003	reanman Smallwood et al
2003/0120170	Al	6/2003	Zhu et al.
2003/0120182	Al	6/2003	Wilkinson et al.
2003/0173976	Al	9/2003	Wiegand et al.
2003/0176808	A1	9/2003	Masuo

References Cited (56)

U.S. PATENT DOCUMENTS

2003/0216661	A1	11/2003	Davies	
2003/0216664	A 1	11/2003	Sucroz	
2005/0210004	AI	11/2003	Suarez	
2004/0015095	Al	1/2004	Li et al.	
2004/0019292	A1	1/2004	Drinan et al.	
2004/0050220	A 1	3/2004	Mourad et al	
2004/0033220	A1	3/2004	Mourad et al.	
2004/0059242	AI	3/2004	Masuo et al.	
2004/0073127	A1	4/2004	Istvan et al.	
2004/0073130	A 1	4/2004	Bohm et al	
2004/0075150	A1	4/2004		
2004/0077944	AI	4/2004	Steinberg et al.	
2004/0116819	A1	6/2004	Alt	
2004/0127703	A 1	7/2004	Mendlein et al	
2004/0127755	A 1 W	0/2004		A CID 5/052
2004/0158167	AI*	8/2004	Smith	A61B 5/053
				600/547
2004/0167422	A 1	8/2004	Dillon at al	000/01/
2004/010/423	AL	0/2004	rmon et al.	
2004/0171961	Al	9/2004	Smith et al.	
2004/0181163	A1	9/2004	Wong et al.	
2004/0181164	A 1	0/2004	Smith at al	
2004/0181104	AI	9/2004	Shilui et al.	
2004/0186392	Al	9/2004	Ward et al.	
2004/0210150	A1	10/2004	Virtanen	
2004/0210158	A 1	10/2004	Organ et al	
2004/0210138	<u></u>	10/2004	Organ et al.	
2004/0220632	AI	11/2004	Burnes	
2004/0234113	A1	11/2004	Miga	
2004/0236202	Δ1	11/2004	Burton	
2004/02/2020	A 1	12/2004	L'anno at al	
2004/0242987	AI	12/2004	Liew et al.	
2004/0242989	A1	12/2004	Zhu et al.	
2004/0243018	Δ1	12/2004	Organ et al	
2004/0243010	A 1	12/2004	Digan et al.	
2004/02528/0	AI	12/2004	Reeves et al.	
2004/0253652	A1	12/2004	Davies	
2004/0260167	A 1	12/2004	Leonhardt	
2004/0200107	A 1	12/2004	K as also as	
2004/0207333	AI	12/2004	Kronberg	
2004/0267344	A1	12/2004	Stett et al.	
2005/0033281	Δ1	2/2005	Bowman et al	
2005/0035201	A 1	2/2005	Karaman et al	
2005/0039763	AI	2/2005	Kraemer et al.	
2005/0049474	A1	3/2005	Kellogg et al.	
2005/0080460	A1	4/2005	Wang et al.	
2005/0085742	A 1	4/2005	Healton at al	
2003/0083743	AI	4/2003	Hacker et al.	
2005/0098343	Al	5/2005	Fukuda	
2005/0101875	A1	5/2005	Semler et al.	
2005/010101075	A 1	5/2005	A set of all	
2005/0107719	AI	5/2005	Arad et al.	
2005/0113704	A1	5/2005	Lawson et al.	
2005/0124908	A1	6/2005	Belalcazar et al.	
2005/0127490	A 1	6/2005	Alt at al	
2005/015/480	AI	0/2005	All et al.	
2005/0151545	Al	7/2005	Park et al.	
2005/0177061	A1	8/2005	Alanen et al.	
2005/0177062	A 1	8/2005	Sizebal at al	
2003/0177002	AI	8/2003	Sklabal et al.	
2005/0192488	Al	9/2005	Bryenton et al.	
2005/0201598	A1	9/2005	Harel et al.	
2005/0203435	A 1	0/2005	Nakada	
2005/0205455	<u>, , , , , , , , , , , , , , , , , , , </u>	0/2005	D	
2005/0203436	AI	9/2005	Davies	
2005/0215918	A1	9/2005	Frantz et al.	
2005/0228309	Δ1	10/2005	Fisher et al	
2005/0220505	A 1	11/2005	$C_{1} = \frac{1}{2} = \frac{1}{2}$	
2005/0251062	AI	11/2005	Choi et al.	
2005/0261743	Al	11/2005	Kroll	
2005/0283091	A1	12/2005	Kink et al.	
2006/0004300	A 1	1/2006	Kennedy	
2000/0004500	4.1	1/2000		
2006/0025701	AI	2/2006	Kasanara	
2006/0041280	A1	2/2006	Stahmann et al.	
2006/0047180	A 1	3/2006	Takehara	
2000/004/109	A 1	3/2000	Dalara	
2006/0052678	AI	3/2006	Drinan	
2006/0064029	A1	3/2006	Arad (Abboud)	
2006/0070623	A 1	4/2006	Wilkinson et al	
2000/00/0025	A 1	4/2006	Course at al	
2000/0085048	AI	4/2006	Cory et al.	
2006/0085049	A1	4/2006	Cory et al.	
2006/0100532	A1	5/2006	Bae et al.	
2006/0111652	A 1	5/2006	McL and	
2000/0111032	AI	5/2000	MCLEOU	
2006/0116599	Al	6/2006	Davis	
2006/0122523	A1	6/2006	Bonmassar et al.	
2006/0122540	A 1	6/2006	Zhu et al	
2000/0122340	AI	0/2000	Znu et al.	
2006/0135886	Al	6/2006	Lippert et al.	
2006/0184060	A1	8/2006	Belalcazar	
2006/0107500	A 1	0/2006	Kanamori et al	
2000/019/309	AI	9/2000	Kanamon et al.	
2006/0200033	A1	9/2006	Keren et al.	
2006/0224079	A1	10/2006	Washchuk	
2000/0224079	4 1	10/2000		
2006/0224080	AI	10/2006	Oku et al.	
2006/0241513	A1	10/2006	Hatlestad et al.	

2006/0241719				
	A1	10/2006	Foster et al.	
2006/0247543	A 1	11/2006	Cornich et al	
2000/0247343	A1	11/2000		
2006/0252670	AI	11/2006	Florucci et al.	
2006/0253016	Al	11/2006	Baker et al.	
2006/0258952	A1	11/2006	Stahmann et al.	
2006/0264775	Δ1	11/2006	Mills et al	
2000/0204775	A 1	11/2006	Stahmann at al	
2000/0204770	AI	11/2006	Stanmann et al.	
2006/0270942	Al	11/2006	Mcadams	
2007/0007975	A1	1/2007	Hawkins et al.	
2007/0010758	A 1	1/2007	Matthiessen et al	
2007/0010750	A 1 *	2/2007	Talmas	CO1D 27/02
2007/0024310	AI ⁺	2/2007	Токипо	GUIK 27/02
				324/610
2007/0027402	A1	2/2007	Levin et al.	
2007/0043303	Δ1	2/2007	Osvoka et al	
2007/0045505	A 1	2/2007	Usfara a st sl	
2007/0049995	AI	5/2007	normann et al.	
2007/0087703	Al	4/2007	Li et al.	
2007/0088227	A1	4/2007	Nishimura	
2007/0106342	A 1	5/2007	Schumann	
2007/0118027	A 1	5/2007	Baltor at al	
2007/0116027	A1	5/2007		
2007/0156061	AI	7/2007	Hess	
2007/0188219	A1	8/2007	Segarra	
2007/0246046	A1	10/2007	Teschner et al.	
2007/0270707	A 1	11/2007	Belalcazar	
2007/0270707	A 1	1/2009	Coolaicazai	
2008/0001608	AI	1/2008	Saumer	
2008/0002873	Al	1/2008	Reeves et al.	
2008/0004904	A1	1/2008	Tran	
2008/0009757	Δ1	1/2008	Tsoglin et al	
2000/0000757	A 1	1/2000	Cl d	
2008/0009759	AI	1/2008	Chetham	
2008/0027350	Al	1/2008	Webler	
2008/0039700	A1	2/2008	Drinan et al.	
2008/0048786	Δ1	2/2008	Feldkamp et al	
2000/0010/00	A 1	2/2008	Pork at al	
2008/0031043	AI	2/2008	Park et al.	
2008/0064981	Al	3/2008	Gregory	
2008/0091114	A1	4/2008	Min et al.	
2008/0139957	A1	6/2008	Hubbard et al.	
2008/0183008	A 1	7/2008	Denison et al	
2008/0183038	A1	0/2008	Demison et al.	
2008/0188/5/	AI	8/2008	Rovira et al.	
2008/0200802	Al	8/2008	Bhavaraju et al.	
2008/0205717	A1	8/2008	Reeves et al.	
2008/0221411	A 1	9/2008	Hausmann et al	
2008/02/111	A 1	10/2008	Lino	
2008/024/302	AI	10/2008		
2008/0252304	AI	10/2008	Woo et al.	
2008/0262375	A1	10/2008	Brown et al.	
2008/0270051	A1	10/2008	Essex et al.	
2008/0287823	A 1	11/2008	Chetham	
2000/0207023		12/2008	T-1-1-	
2008/0306400	AI	12/2008	Takenara	
2008/0306402	Al	12/2008	Singer	
			Word at al	
2008/0319336	Al	12/2008	ward et al.	
2008/0319336 2009/0018432	A1 A1	12/2008	He	
2008/0319336 2009/0018432 2009/0043222	A1 A1	12/2008 1/2009 2/2009	He Chetham	
2008/0319336 2009/0018432 2009/0043222	A1 A1 A1	12/2008 1/2009 2/2009	He Chetham	
2008/0319336 2009/0018432 2009/0043222 2009/0054952	A1 A1 A1 A1	12/2008 1/2009 2/2009 2/2009	He Chetham Glukhovsky et al.	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0069708	A1 A1 A1 A1 A1 A1	12/2008 1/2009 2/2009 2/2009 3/2009	He Chetham Glukhovsky et al. Hatlestad et al.	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0069708 2009/0076343	A1 A1 A1 A1 A1 A1 A1	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009	He Chetham Glukhovsky et al. Hatlestad et al. James et al.	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0069708 2009/0076343 2009/0076345	A1 A1 A1 A1 A1 A1 A1 A1	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009	He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0069708 2009/0076343 2009/0076345	A1 A1 A1 A1 A1 A1 A1 A1	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009	He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al.	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0069708 2009/0076343 2009/0076345 2009/0076350	A1 A1 A1 A1 A1 A1 A1 A1 A1	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009	He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al.	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0076343 2009/0076345 2009/0076350 2009/0076410	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009	He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al.	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0076343 2009/0076345 2009/0076350 2009/0076410 2009/0076410	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009	He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0076343 2009/0076343 2009/0076350 2009/0076410 2009/0084674	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009	He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al.	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0076343 2009/0076343 2009/0076345 2009/0076350 2009/0076410 2009/0082679 2009/0084674	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009	He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl	
2008/0319336 2009/00184322 2009/0043222 2009/0054952 2009/0069708 2009/0076343 2009/0076345 2009/0076410 2009/0082679 2009/0082679 2009/0084674 2009/0093730	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009	He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0069708 2009/0076343 2009/0076345 2009/0076340 2009/0082679 2009/0084674 2009/0084674	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009	He Chetham Glukhovsky et al. Hatlestad et al. James et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al.	
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0076343 2009/0076343 2009/0076345 2009/0076350 2009/0076410 2009/0084674 2009/0093730 2009/0143663	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 6/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham	A61B 5/053
2008/0319336 2009/00184322 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076345 2009/0076410 2009/0082679 2009/0084674 2009/0083730 2009/0105555 2009/0143663	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 6/2009	Wald et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0069708 2009/0076343 2009/0076350 2009/0076350 2009/0076410 2009/0082679 2009/0084674 2009/0084674 2009/0084674 2009/00846555 2009/0143663 2009/0177099	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 6/2009 7/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al.	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076350 2009/0076350 2009/0076350 2009/0084674 2009/0093730 2009/012555 2009/0143663 2009/0177099 2000/020828	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 6/2009 8/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076345 2009/0076410 2009/0076410 2009/0082679 2009/0076410 2009/0082679 2009/016555 2009/0143663 2009/0177099 2009/0209828	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 6/2009 7/2009 8/2009	Wald et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Dacso	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0069708 2009/0076343 2009/0076350 2009/0076350 2009/0082679 2009/0084674 2009/0084674 2009/0084674 2009/0105555 2009/0143663 2009/0177099 2009/0209828 2009/0209872	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 6/2009 7/2009 8/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076350 2009/0076350 2009/0084674 2009/0084674 2009/0093730 2009/017555 2009/0143663 2009/0177099 2009/0209828 2009/0209872 2009/0209872	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 6/2009 8/2009 8/2009 8/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076345 2009/0076345 2009/0082679 2009/0082679 2009/0082679 2009/0184674 2009/0105555 2009/0143663 2009/0177099 2009/0209828 2009/0209828 2009/0216140	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 6/2009 8/2009 8/2009 8/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al.	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0054952 2009/0069708 2009/0076343 2009/0076345 2009/0076350 2009/0084674 2009/0084674 2009/0084674 2009/0084674 2009/017555 2009/0143663 2009/0177099 2009/0209828 2009/0209828 2009/0209872 2009/0216140 2009/0234244	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 8/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076350 2009/0076350 2009/0076350 2009/0084674 2009/0093730 2009/017555 2009/0143663 2009/0177099 2009/0209828 2009/0209872 2009/0209872 2009/0209872 2009/0216140 2009/0216148 2009/0234244	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 6/2009 8/2009 8/2009 8/2009 8/2009 8/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Wobler	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076345 2009/0076350 2009/0076410 2009/0082679 2009/0082679 2009/0082679 2009/0083674 2009/0177099 2009/0177099 2009/0209872 2009/0216140 2009/0216148 2009/0216148 2009/0216148	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 9/2009 9/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076345 2009/0076350 2009/007855 2009/0084674 2009/0084674 2009/0084674 2009/017555 2009/0143663 2009/0177099 2009/0209828 2009/0209828 2009/0209828 2009/0209828 2009/0209828 2009/0209828 2009/0216140 2009/0240163 2009/0240163 2009/0240163	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 8/2009 8/2009 9/2009 10/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz	A61B 5/053 600/372
2008/0319336 2009/00184322 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076345 2009/0076410 2009/0082679 2009/0084674 2009/0093730 2009/0143663 2009/0143663 2009/0177099 2009/0209828 2009/0216140 2009/0216148 2009/0216148 2009/0216148 2009/0244725	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 6/2009 8/2009 8/2009 8/2009 8/2009 8/2009 9/2009 10/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz Markowitz et al.	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076345 2009/0076350 2009/0076410 2009/0082679 2009/0082679 2009/0082679 2009/017099 2009/0177099 2009/0209872 2009/0216140 2009/0216148 2009/0216148 2009/0216148 2009/0240163 2009/0264775	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 8/2009 9/2009 9/2009 9/2009 10/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz et al. Vardy	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076345 2009/0076350 2009/0084674 2009/0084674 2009/0084674 2009/0177099 2009/0177099 2009/0209828 2009/0209828 2009/0209828 2009/0209828 2009/0209828 2009/0216140 2009/0216140 2009/0240163 2009/024772	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 8/2009 8/2009 9/2009 10/2009 10/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz et al. Vardy	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076345 2009/0076345 2009/0076410 2009/0082679 2009/0084674 2009/0084674 2009/0177099 2009/0143663 2009/0177099 2009/0209828 2009/0216140 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/024775 2009/0264775 2009/0264775	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 6/2009 8/2009 8/2009 8/2009 8/2009 8/2009 8/2009 9/2009 10/2009 10/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz et al. Vardy Gregory et al.	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076345 2009/0076350 2009/0076410 2009/0082679 2009/0082679 2009/0082679 2009/0105555 2009/0143663 2009/0177099 2009/0209828 2009/0209872 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/0264777 2009/0264776 2009/0264771 2009/0264791 2009/0264791	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 8/2009 9/2009 9/2009 9/2009 10/2009 10/2009 10/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz et al. Vardy Gregory et al. Zielinski et al.	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076345 2009/0076350 2009/0076350 2009/0084674 2009/0084674 2009/0084674 2009/0177099 2009/0177099 2009/0209828 2009/0209828 2009/0209828 2009/0209828 2009/0209827 2009/0216140 2009/0240163 2009/0240163 2009/0240163 2009/024075 2009/02475	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 8/2009 8/2009 9/2009 10/2009 10/2009 10/2009 11/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz et al. Vardy Gregory et al. Zielinski et al.	A61B 5/053 600/372
2008/0319336 2009/00184322 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076345 2009/0076410 2009/0082679 2009/0084674 2009/0084674 2009/0177099 2009/0143663 2009/0177099 2009/0209828 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/024775 2009/0264776 2009/0264776 2009/02647791 2009/0275854 2009/0275854 2009/0275854	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 6/2009 8/2009 8/2009 8/2009 8/2009 8/2009 9/2009 10/2009 10/2009 10/2009 10/2009 11/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz et al. Vardy Gregory et al. Zielinski et al. Zielinski et al.	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076345 2009/0076350 2009/0076410 2009/0082679 2009/0082679 2009/0082679 2009/0105555 2009/0143663 2009/0177099 2009/0209828 2009/0209872 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/0264777 2009/0264776 2009/0264771 2009/0264755 2009/0275855 2009/0275855	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 9/2009 9/2009 9/2009 9/2009 9/2009 9/2009 10/2009 10/2009 10/2009 11/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz et al. Vardy Gregory et al. Zielinski et al. Zielinski et al. Ward	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076345 2009/0076350 2009/0076350 2009/0084674 2009/0084674 2009/0084674 2009/0177099 2009/0177099 2009/0177099 2009/0209828 2009/0209828 2009/0209828 2009/0209872 2009/0216140 2009/0240163 2009/0240163 2009/024076 2009/02475 2009/024755 2009/0275854 2009/0275855 2009/0275855	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 8/2009 8/2009 9/2009 10/2009 10/2009 10/2009 11/2009 11/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz et al. Vardy Gregory et al. Zielinski et al. Zielinski et al. Ward Davies et al.	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076343 2009/0076345 2009/0076350 2009/0076350 2009/0082679 2009/0082679 2009/0082679 2009/0177099 2009/0177099 2009/0209828 2009/0209828 2009/0209828 2009/0209828 2009/0216140 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/024776 2009/0264776 2009/0264771 2009/0264775 2009/0264775 2009/0264771 2009/0275855 2009/0275854 2009/0275855 2009/027635 2009/027635	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 2/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 8/2009 9/2009 10/2009 10/2009 10/2009 10/2009 11/2009 11/2009 11/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz et al. Vardy Gregory et al. Zielinski et al. Ward Dacso et al.	A61B 5/053 600/372
2008/0319336 2009/0018432 2009/0043222 2009/0069708 2009/0076343 2009/0076345 2009/0076350 2009/0076410 2009/0082679 2009/0082679 2009/0082679 2009/0105555 2009/0143663 2009/0177099 2009/0209872 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/0216148 2009/0264777 2009/0264776 2009/0264776 2009/0264771 2009/0264775 2009/0264775 2009/0264755 2009/0275855 2009/0275855 2009/0287102 2009/0318778	A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A1 A	12/2008 1/2009 2/2009 3/2009 3/2009 3/2009 3/2009 3/2009 4/2009 4/2009 4/2009 4/2009 4/2009 8/2009 8/2009 8/2009 9/2009 9/2009 9/2009 9/2009 9/2009 10/2009 10/2009 11/2009 11/2009 11/2009 12/2009	Ward et al. He Chetham Glukhovsky et al. Hatlestad et al. James et al. Manicka et al. Bly et al. Libbus et al. Chetham Holzhacker et al. Grassl Dacso et al. Chetham Smith et al. Musin Pop Skrabal Freed et al. Tanaka Webler Markowitz et al. Vardy Gregory et al. Zielinski et al. Zielinski et al. Ward Davies et al. Dacso et al. Zielinski et al. Markowitz et al. Zielinski et al.	A61B 5/053 600/372

U.S. PATENT DOCUMENTS

2010/0007357	A1	1/2010	Ammari et al.
2010/0049077	A1	2/2010	Sadleir et al.
2010/0056881	A1	3/2010	Libbus et al.
2010/0094160	A1	4/2010	Eror et al.
2010/0100003	A1	4/2010	Chetham et al.
2010/0100146	A1	4/2010	Blomqvist
2010/0106046	A1	4/2010	Shochat et al.
2010/0152605	A1	6/2010	Ward
2010/0168530	A1	7/2010	Chetham et al.
2010/0191141	A1	7/2010	Aberg
2010/0228143	A1	9/2010	Teschner et al.
2011/0025348	A1	2/2011	Chetham et al.
2011/0034806	A1	2/2011	Hartov et al.
2011/0046505	A1	2/2011	Cornish et al.
2011/0054343	A1	3/2011	Chetham et al.
2011/0054344	A1	3/2011	Slizynski
2011/0060239	A1	3/2011	Gaw
2011/0060241	A1	3/2011	Martinsen et al.
2011/0082383	A1	4/2011	Cory et al.
2011/0087129	A1	4/2011	Chetham et al.
2011/0118619	A1	5/2011	Burton et al.
2011/0190655	A1	8/2011	Moissl et al.
2011/0208084	A1	8/2011	Martinez et al.
2011/0230784	A2	9/2011	Slizynski et al.
2011/0245712	A1	10/2011	Patterson et al.
2011/0251513	A1	10/2011	Chetham
2011/0274327	A1	11/2011	Wehnes et al.
2011/0282180	A1	11/2011	Goldkuhl et al.
2012/0071772	A1	3/2012	Chetham
2012/0165884	A1	6/2012	Xi
2012/0238896	A1	9/2012	Garber et al.
2013/0102873	A1	4/2013	Hamaguchi
2013/0165760	A1	6/2013	Erlinger et al.
2013/0165761	A1	6/2013	De Limon et al.
2014/0148721	A1	5/2014	Erlinger et al.

FOREIGN PATENT DOCUMENTS

~ .				wo
CA	2613524	1/2007		WO
CA	2615845	1/2007		WO
CA	WO 2008119166	A1 * 10/2008	H03F 1/14	WO
CN	1180513	5/1998		WO
CN	1236597	12/1999		WO
CN	1329875	1/2002		WO
DE	2912349	10/1980		WO
EP	0249823	12/1987		WO
EP	349043	3/1990		WO
EP	0357309	3/1990		WO
EP	377887	7/1990		WO
EP	581073	2/1994		WO
EP	339471	3/1997		WO
EP	865763	9/1998		WO
EP	0869360	10/1998		WO
EP	1078597	2/2001		WO
EP	1080686	3/2001		WO
EP	1112715	4/2001		WO
EP	1112715	7/2001		WO
\mathbf{EP}	1146344	10/2001		WO
\mathbf{EP}	1114610	11/2001		WO
EP	1177760	2/2002		WO
\mathbf{EP}	1219937	7/2002		WO
\mathbf{EP}	1238630	9/2002		WO
EP	1247487	10/2002		WO
\mathbf{EP}	1329190	7/2003		WO
\mathbf{EP}	1338246	8/2003		WO
\mathbf{EP}	1080686	3/2004		WO
\mathbf{EP}	1452131	9/2004		WO
EP	1553871	7/2005		WO
\mathbf{EP}	1118308	11/2005		WO
\mathbf{EP}	1629772	3/2006		wo
EP	1247487	1/2008		wo
\mathbf{EP}	1903938	4/2008		wo
EP	1909642	4/2008		wo
EP	1948017	7/2008		WO
EP	1353595	8/2008		WO

FR	2486386	1/1982
ED	2748028	11/1007
GD	2740920	11/1997
GB	1441622	7/19/6
GB	2131558	6/1984
GB	2260416	4/1003
CD	2200410	12/2007
GB	2426824	12/2006
JP	6-74103	10/1994
IP	8191808	7/1006
JI ID	00051004	2/1007
JP	09051884	2/1997
JP	9220209	8/1997
IP	10000185	1/1008
JI ID	10014000	1/1000
JP	10014898	1/1998
JP	10014899	2/1998
JP	10225521	8/1998
ID	11070000	3/1000
JI ID	11070090	3/1333
JP	11-513592	11/1999
JP	2000107138	4/2000
IP	2000139867	5/2000
TD ID	2001027725	2/2001
JP	2001037733	2/2001
JP	2001061804	3/2001
JP	2001-204707	7/2001
ID	2001224568	8/2001
JI ID	2001224508	0/2001
JP	2001-245866	9/2001
JP	2001321352	11/2001
JP	2002502274	1/2002
īD	2002202274	0/2002
JF	2002238870	0/2002
JP	2002330938	11/2002
JP	2002350477	12/2002
TD	2003 502002	1/2003
JI ID	2003-302092	1/2003
JP	2003075487	3/2003
JP	2003-116803	4/2003
JP	2003116805	4/2003
IP	2003230547	8/2003
JI ID	2003250547	2/2003
JP	200401231	2/2004
JP	2006-501892	1/2006
JP	2008-502382	1/2008
IP	2008022995	7/2008
ID	2010 526604	8/2010
	2010-520004	6/2010
KU	2112416	6/1998
WO	WO 88-07392	10/1988
WO	WO 91-19454	12/1991
WO	WO 93-18821	9/1993
WO	WO 04/01040	1/1004
WO	WO 94/01040	1/1994
wo	WO 94-10922	5/1994
WO	WO 96-01586	1/1996
WO	WO 96-12439	5/1996
wo	WO 06 32652	10/1006
WO	WO 90-32032	10/1990
WO	WO 97-11638	4/1997
WO	WO 97-14358	4/1997
WO	WO 97-24156	7/1997
WO	WO 08 06328	2/1008
WO	WO 98-00328	4/1000
WO	WU 98/12983	4/1998
WO	WO 98-23204	6/1998
WO	WO 98-33553	8/1998
WO	WO 98-51211	11/1998
wo	WO 00 42024	0/1000
WO	WO 99-42034	0/1999
wo	WO 99-48422	9/1999
WO	WO 00-19886	4/2000
WO	WO 00-40955	7/2000
WO	WO 00 78212	12/2000
wo	WO 00-78213	12/2000
wO	WO 00-79255	12/2000
WO	WO 01-27605	4/2001
WO	WO 01-50954	7/2001
WÕ	WO 01-52733	7/2001
WO	WO 01-32733	0/2001
WO	WO 01-67098	9/2001
WO	WO 02-053028	7/2002
WO	WO 02-062214	8/2002
WO	WO 02-094096	11/2002
we	WO 04 000115	12/2002
wU	w0 04-000115	12/2003
WO	WO 2004/002301	1/2004
WO	WO 2004/006660	1/2004
wo	WO 2004 021990	2/2004
wO	WO 2004-021880	5/2004
WO	WO 2004-026136	4/2004
WO	WO 2004-030535	4/2004
wo	WO 2004 032739	4/2004
WO	WO 2004-032/38	4/2004
wo	WO 2004-043252	5/2004
WO	WO 2004-047635	6/2004
WO	WO 2004-047636	6/2004
WO	WO 2004-047/030	6/2004
wu	WU 2004-047638	0/2004

FOREIGN PATENT DOCUMENTS

WO	WO 2004-049936	6/2004
WO	WO 2004-083804	9/2004
WO	WO 2004-084087	9/2004
WÕ	WO 2004-084723	10/2004
WÕ	WO 2004-098389	11/2004
wo	WO 2004/112563	12/2004
wõ	WO 2005-010640	2/2005
wo	WO 2005-018432	3/2005
wo	WO 2005-010452	3/2005
wo	WO 2005-027717	6/2005
wo	WO 2005-051105	6/2005
wo	WO 2005-051194	12/2005
wo	WO 2005-122881 WO 2005-122888	12/2005
wo	WO 2005-122888	4/2005
WO	WO 2006-043031	6/2006
wo	WO 2000-030074	12/2006
WO	WO 2006-129108	12/2006
WO	WO 2000-129110 WO 2007 002001	1/2007
WO	WO 2007-002991	1/2007
WO	WO 2007-002992	1/2007
WO	WO 2007-002993	1/2007
WO	WO 2007-009185	1/2007
WO	WO 2007-041785	4/2007
WO	WO 2007/045006	4/2007
WO	WO 2007-030493	5/2007
WO	WO 2007-070997	0/2007
WO	WO 2007/105996	9/2007
WO	WO 2007-128952	1/2007
WO	WO 2008-011/16	1/2008
WO	WO 2008-054426	8/2008
WO	WO 2008/119166	10/2008
WO	WO 2008-138062	11/2008
WO	WO 2008/149125	12/2008
WO.	WO 2009-018620	2/2009
WO	WO 2009-02/812	3/2009
WO	WO 2009-036369	3/2009
WO	WO 2009-068961	6/2009
WO	WO 2009/100491	8/2009
WO	WO 2009-112965	9/2009
WO	WO 2010-003162	1/2010
WO	WO 2010-029465	3/2010
WO	WO 2010-069023	6/2010
WO	WO 2010-076719	7/2010
WO	WO 2011-018744	2/2011
WO	WO 2011-022068	2/2011
wo	WO 2011-050393	5/2011
WO	WO 2011-075769	6/2011
WO	WO 2011-113169	9/2011
ŴΟ	WO 2011-136867	11/2011

OTHER PUBLICATIONS

Al-Hatib, F.; Patient Instrument connection errors in bioelectrical impedance measurement; Physiological Measurement; vol. 19, No. 2, pp. 285-296; May 2, 1998.

Bella, et al., Relations of Left Ventricular Mass to Fat-Free and Adipose Body Mass: The Strong Heart Study, (1998) Circulation, vol. 98, pp. 2538-2544.

Boulier, A. et al.; Fat-Free Mass Estimation by Two Electrode Impedance Method; American Journal of Clinical Nutrition; vol. 52, pp. 581-585; 1990.

Bracco, D. et al., Bedside determination of fluid accumulation after cardiac surgery using segmental bioelectrical impedance, Critical Care Medicine, vol. 26, No. 6, pp. 1065-1070, 1998.

Chaudary, S.S. et al.; Dielectric Properties of Normal & Malignant Human Breast Tissues at Radiowave and Microwave Frequencies; Indian Journal of Biochemistry & Biophysics; vol. 21, No. 1, pp. 76-79; 1984.

Chiolero, R.L. et al.; Assessment of changes in body water by bioimpedance in acutely ill surgical patients; Intensive Care Medicine; vol. 18, pp. 322-326; 1992.

Chumlea et al.; Bioelectrical Impedance and Body Composition: Present Status and Future Directions; Nutrition Reviews; vol. 52, No. 4, pp. 123-131; 1994. Cornish, B.H. et al.; Alteration of the extracellular and total body water volumes measured by multiple frequency bioelectrical impedance analysis; Nutrition Research; vol. 14, No. 5, pp. 717-727; 1994.

Cornish, B.H. et al.; Bioelectrical impedance for monitoring the efficacy of lymphoedema treatment programmes; Breast Cancer Research and Treatment; vol. 38, pp. 169-176; 1996.

Cornish, B.H. et al.; Data analysis in multiple-frequency bioelectrical impedance analysis; Physiological Measurement; vol. 19, No. 2, pp. 275-283; May 1, 1998.

Cornish, B.H. et al.; Early diagnosis of lymphedema using multiple frequency bioimpedance; Lymphology; vol. 34, pp. 2-11; Mar. 2001.

Cornish, B.H. et al.; Early diagnosis of lymphoedema in postsurgery breast cancer patients; Annals New York Academy of Sciences; pp. 571-575; May 2000.

Cornish, B.H. et al.; Quantification of Lymphoedema using Multifrequency Bioimpedance; Applied Radiation and Isotopes; vol. 49, No. 5/6, pp. 651-652; 1998.

De Luca, F. et al., Use of low-frequency electrical impedance measurements to determine phospoholipid content in amniotic fluid; Physics in Medicine and Biology, vol. 41, pp. 1863-1869, 1996.

Deurenberg, P. et al., Multi-frequency bioelectrical impedance: a comparison between the Cole-Cole modelling and Hanai equations with the classically impedance index approach, Annals of Human Biology, vol. 23, No. 1, pp. 31-40, 1996.

Dines K.A. et al.; Analysis of electrical conductivity imaging; Geophysics; vol. 46, No. 7, pp. 1025-1036; Jul. 1981.

Ellis, K.J. et al; Human hydrometry: comparison of multifrequency bioelectrical impedance with 2H2O and bromine dilution; Journal of Applied Physiology; vol. 85, No. 3, pp. 1056-1062; 1998.

Ezenwa, B.N. et al.; Multiple Frequency System for Body Composition Measurement; Proceedings of the Annual International Conference of the Engineering in Medicine and Biology Society; vol. 15; pp. 1020-1021; 1993.

Forslund, A.H. et al.; Evaluation of modified multicompartment models to calculate body composition in healthy males; American Journal of Clinical Nutrition; vol. 63, pp. 856-862; 1996.

Gersing, E.; Impedance spectroscopy on living tissue for determination of the state of Organs; Bioelectrochemistry and Bioenergetics; vol. 45, pp. 145-149; 1998.

Gerth, W.A. et al.; A computer-based bioelectrical impedance spectroscopic system for noninvasive assessment of compartmental fluid redistribution; Third Annual IEEE Symposium on Computer Based Medical Systems, Jun. 3-6, 1990, University of NC. At Chapel Hill; pp. 446-453; Jun. 1990.

Gudivaka R. et al; Single- and multifrequency models for bioelectrical impedance analysis of body water compartments; Applied Physiology; vol. 87, Issue 3, pp. 1087-1096; 1999.

Iacobellis, G., et al. Influence of Excess Fat on Cardiac Morphology and Function: Study in Uncomplicated Obesity, (2002) Obesity Research, vol. 10, pp. 767-773.

Ivorra, A., et al.; Bioimpedance Dispersion Width as a Parameter to Monitor Living Tissues; Physiological Measurement; vol. 26; pp. 1-9; 2005.

Jones, C.H. et al; Extracellular fluid volume determined by bioelectric impedance and serum albumin in CAPD patients; Nephrology Dialysis Transplantation; vol. 13, pp. 393-397; 1998. Jossinet, J. et al.; A Study for Breast Imaging with a Circular Array

of Impedance Electrodes; Proc. Vth Int. Conf. Bioelectrical Impedance, 1981, Tokyo, Japan; pp. 83-86; 1981.

Jossinet, J. et al.; Technical Implementation and Evaluation of a Bioelectrical Breast Scanner; Proc. 10.sup.th Int. Conf. IEEE Engng. Med. Biol., 1988, New Orleans, USA (Imped. Imaging II); vol. 1. p. 289; 1988.

Kanai, H. et al.; Electrical Measurement of Fluid Distribution in Legs and Arms; Medical Progress through technology; pp. 159-170; 1987.

Karason, K., et al., Impact of Blood Pressure and Insulin on the Relationship Between Body Fat and Left Ventricular Structure, (2003) European Heart Journal, vol. 24, pp. 1500-1505.

OTHER PUBLICATIONS

Kim, C.T. et al.; Bioelectrical impedance changes in regional extracellular fluid alterations; Electromyography and Clinical Neurophysiology; vol. 37, pp. 297-304; 1997.

Liu R. et al; Primary Multi-frequency Data Analyze in Electrical Impedance Scanning; Proceedings of the IEEE-EMBS 2005, 27th Annual International Conference of the Engineering in Medicine and Biology Society, Shanghai, China; pp. 1504-1507; , Sep. 1-4, 2005.

Lozano, A. et al.; Two-frequency impedance plethysmograph: real and imaginary parts; Medical & Biological Engineering & Computing; vol. 28, No. 1, pp. 38-42; Jan. 1990.

Lukaski, H.C. et al.; Estimation of Body Fluid Volumes Using Tetrapolar Bioelectrical Impedance Measurements; Aviation, Space, and Environmental Medicine; pp. 1163-1169; Dec. 1988.

Man, B. et al. Results of Preclinical Tests for Breast Cancer Detection by Dielectric Measurements; XII Int. Conf. Med. Biol. Engng. 1979, Jerusalem, Israel. Springer Int., Berlin; Section 30.4; 1980.

Mattar, J.A., Application of Total Body Impedance to the Critically Ill Patient, New Horizons, vol. 4, No. 4, pp. 493-503, Nov. 1996. McCullah, et al.; Bioelectrical Impedance Analysis Measures the Ejection Fraction of the Calf Muscle Pump; IFMBE Proceedings; vol. 17, pp. 616-619; 2007.

McDougal D., et al.; Body Composition Measurements From Whole Body Resistance and Reactance; Surgical Forum; vol. 36, pp. 43-44; 1986.

Osterman K.S. et al.; Multifrequency electrical impedance imaging: preliminary in vivo experience in breast; Physiological Measurement; vol. 21, No. 1, pp. 99-109; Feb. 2000.

Ott, M. et al.; Bioelectrical Impedance Analysis as a Predictor of Survival in Patients with Human Immunodeficiency Virus Infection; Journal of Acquired Immune Deficiency Syndromes and Human Retrovirology; vol. 9, pp. 20-25; 1995.

Pethig, R. et al.; The Passive Electrical Properties of Biological Systems: Their Significance in Physiology, Biophysics and Biotechnology; Physics in Medicine and Biology; vol. 32, pp. 933-970; 1987.

Piperno, G. et al.; Breast Cancer Screening by Impedance Measurements; Frontiers of Medical & Biological Engineering; vol. 2, pp. 111-117; 1990.

Rigaud, B. et al.; Bioelectrical Impedance Techniques in Medicine; Critical Reviews in Biomedical Engineering; vol. 24 (4-6), pp. 257-351; 1996.

Scharfetter, H. et al.; Effect of Postural Changes on the Reliability of Volume Estimations from Bioimpedance Spectroscopy Data; Kidney International; vol. 51, No. 4, pp. 1078-2087; 1997.

Schneider, I.; Broadband signals for electrical impedance measurements for long bone fractures; Engineering in Medicine and Biology Society, 1996. Bridging Disciplines for Biomedicine. Proceedings of the 18th Annual International Conference of the IEEE; vol. 5, pp. 1934-1935; Oct. 31, 1996.

Skidmore, R. et al.; A Data Collection System for Gathering Electrical Impedance Measurements from the Human Breast; Clinical Physics Physiological Measurement; vol. 8, pp. 99-102; 1987. Sollish, B.D. et al.; Microprocessor-assisted Screening Techniques; Israel Journal of Medical Sciences; vol. 17, pp. 859-864; 1981.

Steijaert, M. et al.; The use of multi-frequency impedance to determine total body water and extracellular water in obese and lean female individuals; International Journal of Obesity; vol. 21, pp. 930-934; 1997.

Surowiec, A.J. et al.; Dielectric Properties of Brest Carcinoma and the Surrounding Tissues; IEEE Transactions on Biomedical Engineering; vol. 35, pp. 257-263; 1988.

Tedner, B.; Equipment Using Impedance Technique for Automatic Recording of Fluid-Volume Changes During Haemodialysis; Medical & Biological Engineering & Computing; pp. 285-290; 1983.

Thomas. B.J. et al.; Bioelectrical impedance analysis for measurement of body fluid volumes—A review; Journal of Clinical Engineering; vol. 17, No. 16, pp. 505-510; 1992. Thomas. B.J. et al.; Bioimpedance Spectrometry in Determination of Body Water Compartments: Accuracy and Clinical Significance; Applied Radiation and Isotopes; vol. 49, No. 5/6, pp. 447-455; 1998.

Thomas. B.J.; Future Technologies; Asia Pacific Journal Clinical Nutrition; vol. 4, pp. 157-159; 1995.

Ulgen, Y. et al.; Electrical parameters of human blood; Engineering in Medicine and Biology Society, 1998. Proceedings of the 20th Annual International Conference of the IEEE; vol. 6, pp. 2983-2986; Nov. 1, 1998.

Ward, L.C. et al., Multi-frequency bioelectrical impedance augments the diagnosis and management of lymphoedema in postmastectomy patients, European Journal of Clinical Investigation, vol. 22, pp. 751-754, 1992.

Ward, L.C. et al.; Determination of Cole parameters in multiple frequency bioelectrical impedance analysis using only the measurement of impedances; Four-frequency fitting; Physiological Measurement; vol. 27, No. 9, pp. 839-850; Sep. 2006.

Ward, L.C. et al.; There is a better way to measure Lymphoedema; National Lymphedema Network Newsletter; vol. 7, No. 4, pp. 89-92; Oct. 1995.

Woodrow, G. et al; Effects of icodextrin in automated peritoneal dialysis on blood pressure and bioelectrical impedance analysis; Nephrology Dialysis Transplantation; vol. 15, pp. 862-866; 2000. Yamakoshi, K.; Non-Invasive Cardiovascular Hemodynamic Measurements; Sensors in Medicine and Health Care; pp. 107-160; 2004.

Yoshinaga, M., Effect of Total Adipose Weight and Systemic Hypertension on Left Ventricular Mass in Children, American Journal of Cardiology, (1995) vol. 76, pp. 785-787.

International Search Report and Written Opinion of the International Searching Authority issued in PCT/AU2006/000922 dated Oct. 10, 2006.

International Search Report and Written Opinion of the International Searching Authority issued in PCT/AU2008/000588 dated Aug. 13, 2008.

International Search Report from International Application No. PCT/AU2006/000924 dated Sep. 27, 2006.

Cornish, et al., "Optimizing Electrode Sites for Segmental Bioimpedance Measurements" Physiological Measurement, Institute of Physics, 1999, pp. 241-250, vol. 20, No. 3.

Cornish, et al., "A New Technique for the Quantification of Peripheral Edema with Application in Both Unilateral and Bilateral Cases" Angiology, 2002, pp. 41-47, vol. 53, No. 1.

Fenech, et al., "Extracellular and Intracellular Volume Variations During Postural Change Measured by Segmental and Wrist-Ankle Bioimpedance Spectroscopy" IEEE Transactions on Biomedical Engineering, IEEE Service Center, 2004, pp. 166-175, vol. 51, No. 1.

Golden, et al., "Assessment of Peripheral Hemodynmics using Impedance Plethysmogrphy" Physical Therapy, 1986, pp. 1544-1547, vol. 66, No. 10.

Kim, et al., "Impedance Tomography and its Application in Deep Venous Thrombosis Detection" IEEE Engineering in Medicine and Biology Magazine, IEEE Service Center, 1989, pp. 46-49, vol. 8, No. 1.

Nawarycz, et al., "Triple-frequency Electroimpedance Method for Evaluation of Body Water Compartments" Medical & Biological Engineering & Computing, 1996, pp. 181-182, vol. 34, No. Supp. 01, Pt. 02.

Noshiro, et al., "Electrical Impedance in the Lower Limbs of Patients with Duchenne Muscular Dystrophy: A Preliminary Study" Medical & Biological Engineering & Computing, 1993, pp. 97-102, vol. 31, No. 2.

Seo, et al., "Measuring Lower Leg Swelling: Optimum Frequency for Impedance Method" Medical & Biological Engineering & Computing, 2001, pp. 185-189, vol. 39.

Seoane, et al., "Current Source for Wideband Electrical Bioimpedance Spectroscopy Based on a Single Operational Amplifier" World Congress on Medical Physics and Biomedical Engineering 2006, pp. 707-710, vol. 14.

Smith, et al., "A Pilot Study for Tissue Characterization Using Bio-impedance Mapping" 13th International Conference on Elec-

OTHER PUBLICATIONS

trical Bio-impedance and the 8th Conference on Electrical Impedance Tomography 2007, pp. 146-149.

Stanton, et al., "Non-invasive Assessment of the Lymphedematous Limb" Lymphology, The International Society of Lymphology, 2000, pp. 122-135, vol. 33, No. 3.

Bernstein; A New Stroke Volume Equation for Thoracic Electrical Bio Impedance; Critical Care Medicine; vol. 14, pp. 904-909; 1986. Blad et al.; Impedance Spectra of Tumour Tissue in Tomparison with Normal Tissue; A Possible Clinical Application for Electrical Impedance Tomography; Physiological Measurement; vol. 17, pp. A105-A115; 1996.

De Lorenzo et al.; Determination of Intracellular Water by Multifrequency Bioelectrical Impedance; Ann. Nutr. Metab.; vol. 39, pp. 177-184; 1995.

Edwards, L.S.; A Modified Pseudosection for Resistivity and IP; Geophysics; vol. 42, No. 5, pp. 1020-1036; 1977.

Hansen, E.; On the Influence of Shape and Variations in Conductivity of the Sample on Four-Point Measurements; Applied Scientific Research; Section B; vol. 8, Issue 1, pp. 93-104 1960.

Igel, J.; On the Small-Scale Variability of Electrical Soil Properties and Its Influence on Geophysical Measurements; Ph.D. Thesis; Frankfurt University; Hanover, Germany; p. 188; 2007.

Kyle et al.; Bioelectrical Impedance Analysis—Part I: Review of Principles and Methods; Clinical Nutrition; vol. 23, pp. 1226-1243; 2004.

Loke et al.; Least Squares Deconvolution of Apparent Resistivity Pseudosections; Geophysics; vol. 60, No. 6, pp. 1682-1690; 1995. McAdams et al; Tissue Impedance: a Historical Overview Physiological Measurement; Institute of Physics Publishing; vol. 16. (3A), pp. A1-A13; 1995. McEwan et al.; Battery Powered and Wireless Electrical Impedance Tomography Spectroscopy Imaging Ssing Bluetooth; Medicon IFMBE Proceedings; vol. 16, pp. 798-801; 2007.

Roy, A.; Depth of investigation in Direct Current Methods Geophysics; vol. 36, pp. 943-959; 1971.

Wilson et al.; Feasibility Studies of Electrical Impedance Spectroscopy for Monitoring Tissue Response to Photodynamic Therapy; Optical Methods for Tumor Treatment and Detections: Mechanisms and Techniques in Photodynamic Therapy VII; Proc. SPIE 3247; pp. 69-80; 1998.

International Search Report and Written Opinion of the International Searching Authority issued in PCT/AU2009/000163 dated Apr. 16, 2009.

International Search Report and Written Opinion of the International Searching Authority issued in PCT/AU2006/000922 dated Oct. 13, 2006.

International Search Report and Written Opinion of the International Searching Authority issued in PCT/AU2006/001057 dated Oct. 25, 2006.

International Search Report and Written Opinion of the International Searching Authority issued in PCT/AU2008/000034 dated Mar. 17, 2008.

International Search Report and Written Opinion of the International Searching Authority issued in PCT/CA2008/000588 dated Aug. 13, 2008.

International Search Report and Written Opinion of the International Searching Authority issued in PCT/AU2006/000924 dated Oct. 5, 2006.

International Search Report and Written Opinion of the International Searching Authority issued in PCT/AU2008/001521 dated Jan. 15, 2009.

* cited by examiner



Fig. 1



Fig. 2



Fig. 3









Fig. 6





Fig. 8



Fig. 9



SIGNAL DISTRIBUTION FOR PATIENT-ELECTRODE MEASUREMENTS

RELATED APPLICATIONS

This application is a U.S. National Phase under 35 U.S.C. §371 of the International Patent Application No. PCT/ AU2010/001552, filed Nov. 18, 2010, and published in English on May 5, 2011 as WO 2011/060497, which claims the benefit of Australian Patent Application No. 2009905642, filed Nov. 18, 2009, both of which are incorporated by reference in their entirety.

The section headings used herein are for organizational purposes only and are not to be construed as limiting the 15 subject matter described in any way.

BACKGROUND OF THE INVENTION

The present invention relates to apparatus for electrically connecting measurement apparatus to a biological subject, and in particular, to a circuit for electrical impedance measurements, which in one example allows drive signal injection and signal amplitude and phase measurement.

DESCRIPTION OF THE PRIOR ART

The reference in this specification to any prior publication (or information derived from it), or to any matter which is 30 known, is not, and should not be taken as an acknowledgment or admission or any form of suggestion that the prior publication (or information derived from it) or known matter forms part of the common general knowledge in the field of endeavour to which this specification relates. 35

One existing technique for determining biological indicators relating to a subject, such as cardiac function, body composition, and other health status indicators, such as the presence of oedema, involves the use of bioelectrical impedance. This process typically involves using a measuring ⁴⁰ device to measure the electrical impedance of a subject's body using a series of electrodes placed on the skin surface. Changes in electrical impedance measured at the body's surface are used to determine parameters, such as changes in fluid levels, associated with the cardiac cycle, oedema, or the ⁴⁵ like.

Impedance measuring apparatus is sometimes sensitive to external factors, including stray capacitances between the subject and the local environment and the measurement apparatus, variations in electrode/tissue interface impedances, also known as electrode impedances, as well as stray capacitances and inductive coupling between the leads used to connect the measuring device to the electrodes.

It will be appreciated that similar issue also arise when 55 making other electrical measurements relating to biological subjects.

WO2009/059351 describes apparatus for use in performing impedance measurements on a subject. The apparatus includes a processing system for causing a first signal to be 60 applied to the subject, determining an indication of a second signal measured across the subject, using the indication of the second signal to determine any imbalance and if an imbalance exists, determining a modified first signal in accordance with the imbalance and causing the modified 65 first signal to be applied to the subject to thereby allow at least one impedance measurement to be performed.

SUMMARY OF THE PRESENT INVENTION

In a first broad form the present invention provides apparatus for electrically connecting measurement apparatus to a biological subject, the apparatus including a signal delivery circuit including:

a) a current buffer having:

- i) a current buffer input for receiving a signal from a signal source; and,
- ii) a current buffer output for supplying a current to an electrode attached to the biological subject; and,
- b) a voltage buffer having:
 - i) a voltage buffer input coupled to the current buffer output; and,
 - ii) a voltage buffer output for providing a voltage signal indicative of a voltage at the electrode, to a sensor.

Typically the current buffer is a current conveyor.

Typically the voltage buffer is an amplifier connected as ₂₀ an output follower.

Typically the apparatus includes an offset control circuit coupled between the voltage buffer output and the current buffer input.

Typically the offset control circuit includes an integrator coupled between the voltage buffer output and the current buffer input.

Typically the offset control circuit is used to control a DC offset at the electrode.

Typically the signal delivery circuit includes a negative impedance circuit coupled between the voltage buffer output and the current buffer output.

Typically the negative impedance circuit includes a negative impedance circuit amplifier and a compensation impedance.

Typically at least one of a gain of the negative impedance circuit amplifier and a value of the compensation impedance is selected to compensate for parasitic impedance losses.

- Typically the negative impedance circuit includes:
- a) a compensation impedance having a compensation impedance value, a first terminal and a second terminal, the first terminal coupled to the current buffer output; and,
- b) a compensation amplifier having a compensation amplifier input, a compensation amplifier output and a gain, the compensation amplifier input coupled to the voltage buffer output, the compensation amplifier output coupled to the second terminal of the compensation impedance to provide a compensation current to flow through the impedance, and the gain and the compensation impedance values are selected based on the parasitic impedance value so that the compensation current has a magnitude substantially equal to the leakage current magnitude.

Typically the apparatus includes:

a) a plurality of signal delivery circuits, each signal delivery circuit being for supplying a current to a respective electrode attached to the biological subject;
 b) at least one signal source; and

- b) at least one signal source; and,
- c) at least one first multiplexer for selectively connecting the signal source to one of the plurality of signal delivery circuits to thereby apply a current to the biological circuit via the respective electrode.
- Typically the apparatus includes:
- a) a plurality of signal delivery circuits, each signal delivery circuit being for providing a voltage signal indicative of a voltage at the electrode, to a sensor;
- b) at least one sensor; and,

c) at least one second multiplexer for selectively connecting the at least one sensor to one of the plurality of signal delivery circuits to thereby allow the sensor to sense the voltage signal indicative of a voltage at the respective electrode.

Typically the apparatus includes, first and second signal sources for generating respective first and second drive signals, the first and second signal sources being coupled to respective signal delivery circuits.

Typically the first and second signal sources are for 10 generating at least one of:

a) complementary signals;

b) signals having a constant amplitude and phase; and,

c) signals having a controlled amplitude and phase.

Typically at least one signal source is coupled to ground. 15

- Typically the apparatus includes:
- a) at least one signal delivery circuit for supplying a current to a drive electrode attached to the biological subject; and,
- b) at least one signal delivery circuit for providing a 20 voltage signal indicative of a voltage at a sense electrode.

Typically the measurement apparatus includes at least one signal source and at least one sensor.

Typically the measurement apparatus is an impedance 25 measurement apparatus.

In a second broad form the present invention provides a current multiplexing system comprising:

- a) a plurality of current multiplexers having at least one input and a plurality of outputs;
- b) a first and second alternating-current current source each current source having a constant magnitude and frequency, wherein the second current source is complimentary to the first current source, each of the first and second current sources is coupled to an input of a current multiplexer;
- c) a plurality of current delivery circuits, each current delivery circuit having an input, a current output and a voltage output, wherein each current delivery circuit input is coupled to an output of a current multiplexer; 40
- d) a plurality of electrodes, each of the plurality of electrodes being coupled to a current output of one of the plurality of current delivery circuits; and
- e) at least one voltage multiplexer having a plurality of inputs and at least one output;

f) wherein each current delivery circuit comprises:

- i) a current conveyor, having an input and an output, the current conveyor input being coupled to the current delivery circuit input and the current conveyor output being coupled to the current delivery circuit 50 current output;
- ii) a voltage buffer, having an input and an output, the voltage buffer input being coupled to the current delivery circuit current output and the voltage buffer output being coupled to the current delivery circuit 55 voltage output;
- iii) an integrator coupled between the voltage buffer output and the current conveyor input; and,
- iv) a negative impedance circuit coupled between the voltage buffer output and the current conveyor out- 60 put, the negative impedance comprising:
 - (1) a compensation impedance having a compensation impedance value, a first terminal and a second terminal, the first terminal coupled to the current conveyor output;
 - (2) a compensation amplifier having a compensation amplifier input, a compensation amplifier output

4

and a gain, the compensation amplifier input coupled to the voltage buffer output, the compensation amplifier output coupled to the second terminal of the compensation impedance to provide a compensation current to flow through the impedance, and the gain and the compensation impedance values are selected based on the parasitic impedance value so that the compensation current has a magnitude substantially equal to the leakage current magnitude.

In a third broad form the present invention provides apparatus for electrically connecting measurement apparatus to a biological subject, the apparatus including a signal delivery circuit including:

a) a first buffer having:

- i) a first buffer input for receiving a signal from a signal source; and,
- ii) a first buffer output for supplying a drive signal to an electrode attached to the biological subject; and,
- b) a second buffer having:
 - i) a second buffer input coupled to the first buffer output; and,
 - ii) a second buffer output for providing a sensed signal indicative of a signal at the electrode, to a sensor.

Typically the first buffer is a current buffer, the drive signal being a drive current.

Typically the current buffer is a current conveyor.

Typically the second buffer is a voltage buffer the sensed signal being a voltage signal indicative of a voltage at the electrode.

Typically the second buffer is a current buffer the sensed signal being indicative of the drive current.

plimentary to the first current source, each of the first and second current sources is coupled to an input of a 35 current multiplexer; In a fourth broad form the present invention provides on a subject, wherein the apparatus includes:

- a) a signal source for applying a signal to a subject;
- b) a sensor for sensing signals from the subject;
- c) a processing system for controlling the signal source and receiving signals from the sensor; and,
- d) at least one signal delivery circuit, including:
 - i) a first buffer having:
 - (1) a first buffer input for receiving a signal from a signal source; and,
 - (2) a first buffer output for supplying a drive signal to an electrode attached to the biological subject; and,
 - ii) a second buffer having:
 - (1) a second buffer input coupled to the first buffer output; and,
 - (2) a second buffer output for providing a sensed signal indicative of a signal at the electrode, to a sensor.

Typically the apparatus includes:

- a) at least one signal delivery circuit for supplying a current to a drive electrode attached to the biological subject; and,
- b) at least one signal delivery circuit for providing a voltage signal indicative of a voltage at a sense electrode.

Typically the apparatus includes:

- a) a first signal delivery circuit coupled to a drive electrode attached to the biological subject, the first signal delivery circuit including:
 - i) a first buffer having:
 - (1) a first buffer input for receiving a signal from the signal source; and,

(2) a first buffer output for supplying the drive signal to the drive electrode; and,

- ii) a second buffer having:
 - (1) a second buffer input coupled to the first buffer output; and,
 - (2) a second buffer output for providing a sensed signal indicative of the drive signal at the drive electrode;
- b) a second signal delivery circuit coupled to a sense electrode attached to the biological subject, the second signal delivery circuit including,
 - i) a first buffer having:
 - (1) a first buffer input coupled to ground; and,
 - (2) a first buffer output coupled to the sense elec- $\frac{15}{15}$ trode; and,
 - ii) a second buffer having:
 - (1) a second buffer input coupled to the first buffer output; and,
 - (2) a second buffer output for providing a sensed 20 signal indicative of a sensed voltage at the sense electrode.

BRIEF DESCRIPTION OF THE DRAWINGS

The skilled person in the art will understand that the drawings, described below, are for illustration purposes only. The drawings are not intended to limit the scope of the applicants' teachings in any way.

An example of the present invention will now be 30 described with reference to the accompanying drawings, in which:

FIG. 1 is a schematic diagram of a current multiplexing system;

FIG. 2 is a schematic diagram of a current multiplexing ³⁵ system according to various embodiments of applicants' teachings;

FIG. 3 is a schematic diagram of the current multiplexing system of FIG. 2 showing additional components;

FIG. 4 is a schematic diagram of the current multiplexing 40 system of FIG. 2 showing the current delivery circuit in greater detail;

FIG. 5 is a schematic diagram of the current multiplexing system of FIG. 4 showing additional components;

FIG. 6 is a schematic diagram of the negative impedance 45 of FIG. 5.

FIG. 7 is a schematic diagram of the current multiplexing system of FIG. 5 showing the DC bias control and negative impedance in greater detail;

FIG. 8 is a schematic diagram of the negative impedance 50 of FIG. 7;

FIG. 9 is a schematic diagram of an example of an impedance measuring device; and,

FIG. 10 is a flowchart of an example of a process for performing impedance measuring.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Electronic testing equipment or measuring apparatus may 60 be electrically connected with a test subject through electrodes. These electrodes may be used to, for example, deliver currents and measure voltages at various points of contact between the test subject and the electrodes. An example of such test equipment or measuring apparatus can 65 include medical equipment, in which case the test subject can be a biological subject such as a human patient.

6

Regardless of the type of application, a signal source, such as a current source may be used to deliver a drive signal, such as a current, to the electrodes. Often a single current source may be used for multiple electrodes by switching the currents to the various electrodes as needed. FIG. 1 illustrates a current multiplexing system 100. Current multiplexing system 100 comprises a current source 102 coupled to the input of multiplexer 104. Each of the outputs of multiplexer 104 can in turn be coupled to a plurality of electrodes 106. For clarity FIG. 1, only illustrates a single electrode. Each of electrodes 106 can be coupled to a patient 108.

Current source 102 is used to deliver a current I to the input of multiplexer 104. Multiplexer 104 is then used to switch the input current to a particular output and thereby to a particular electrode 106. However, multiplexer 104 can cause a large parasitic impedance 110 to appear at each of the outputs. Consequently, some of the current outputted by multiplexer 104 is lost to parasitic impedance 110 as a leakage current (ΔI). Therefore, only a portion ($I-\Delta I$) of the current produced by current source 102 reaches the electrode 106. The amount of current lost is proportional to the impedance of the interface between electrode 106 and patient 108. Each individual patient may have a different impedance associated with their skin and/or tissue. Consequently, the proportion of the total current that is lost to parasitic impedance 110 may differ with each individual patient. Accordingly, the proportion of total current delivered to the patient may also differ with each individual patient. Thus, with current multiplexing system 100 it can be difficult to deliver a precise amount of current to a patient 108 via an electrode 106.

Reference is now made to FIG. 2, which illustrates various embodiments of current multiplexing system 200, according to various embodiments of applicants' teachings. Current multiplexing system 200 comprises a current source 202, one or more multiplexers 204, a plurality of electrodes 206 coupled to patient 208, one or more voltage multiplexers 212, and a plurality of current delivery circuits 220. Node 214, which is the output of multiplexer 212, can be used to obtain voltage measurements.

Each output of each multiplexer 204 can be connected to a current delivery circuit similar to current delivery circuit 220, which in turn can be connected to the inputs of multiplexer 212. For clarity, FIG. 2 only illustrates a single connection to the output of multiplexer 204 and a single connection to multiplexer 212. The actual number of multiplexers 204, current delivery circuits 220, and multiplexers 212 used depends on such factors as the number electrodes 206 that are coupled to the patient, the number of outputs available on each multiplexer 204, and the number of inputs available on multiplexer 212.

In various embodiments according to applicants teachings, current delivery circuit 220 receives a current I at node 55 222 and outputs a substantially equal current at node 224. In addition, in various embodiments, current delivery circuit 220 produces a voltage at node 226 that is substantially equal to the voltage at node 224. As was explained above, node 226 is coupled to voltage multiplexer 212. However, it should be understood that voltage measurements can be taken directly at node 226, which is the voltage output of delivery circuit 220. The use of voltage multiplexer 212 is optional and can be used to reduce the number of nodes to which voltage measurement equipment is to be connected to.

As will be described in greater detail herein, in various embodiments, current multiplexing system 200 can utilize a single controlled constant current source 202 to deliver a current that is multiplexed to numerous electrical loads, such as electrodes 206. In some embodiments, constant current source 202 delivers an AC current with a constant amplitude and phase, which may for example be, but not limited to, a 5 sinusoid of fixed frequency.

In some embodiments, current multiplexing system 200 can deliver a current to a load that is substantially equal to the current outputted by 202. In various embodiments, current multiplexing system 200 can be utilized to deliver a 10 controllable constant current to numerous locations and loads with a small transmission loss. In addition, in various embodiments, current multiplexing system 200 can be used without a calibration step prior to use. Furthermore, in various embodiments, the loads to which current multiplex- 15 ing system 200 connects need not be altered in any manner in order to utilize current multiplexing system 200. In some embodiments, current delivery circuit 200 can be utilized with any appropriate equipment including but not limited to measuring apparatus such as medical instrumentation, bio- 20 a current injection electrode and electrode 2066 could be metric instrumentation, electrocardiograph equipment, impedance measuring apparatus, and circuit testing equipment

In various embodiments, current delivery circuit 220 can be integrated on a single integrated circuit. In some embodi- 25 ments, a plurality of current delivery circuit, such as delivery circuit 220 can be integrated on a single integrated circuit and packaged in a single chip. In some embodiments, one or more current multiplexers, such as multiplexer 204, a plurality of delivery circuits and one or more voltage multi- 30 plexers, such as multiplexer 212, can be integrated on a single integrated circuit and packaged in a single chip.

Reference is now made to FIG. 3, which illustrates two current delivery circuits 220a and 220b of the current multiplexing system 200 of FIG. 2. As was mentioned above 35 with reference to FIG. 2, in various embodiments, current multiplexing system 200 comprises a plurality of current delivery circuits 220 and a plurality of electrodes. FIG. 2 illustrates only a single current delivery circuit and a single electrode for reasons of clarity. However, as mentioned 40 above, in various embodiments, current multiplexing system 200 comprises at least two current delivery circuits and at least two electrodes, where one electrode can be a current delivery electrode and the second electrode can be a current return electrode. 45

FIG. 3 illustrates two current delivery circuits and two electrodes. It should be understood however that a greater number could be used. Nonetheless, in some embodiments, two current delivery circuits and two electrodes are sufficient. For example, but not limited to, electrode 206a could 50 be used as a current delivery electrode and electrode 206b could be used as a current return electrode. In such embodiments, one of electrodes 206a and 206b could be used as a positive voltage measurement electrode and the other could be used as a negative voltage measurement electrode. The 55 voltage at each of nodes 214a and 214b, which are the outputs of multiplexers 212a and 212b respectively, could be measured to obtain a differential voltage measurement between electrodes 206a and 206b. This may be referred to as a two point measurement given that two electrodes are 60 used (for both current injection/return and for voltage measurement) and therefore there are two points of contact between the current multiplexing system and the patient.

FIG. 3, shows the input of multiplexer 204b connected to ground. In various other embodiments used for a two point 65 measurement, the input of multiplexer can be, for example but not limited to, connected to a second current source (not

shown) that is complimentary to current source 202. More specifically, the second current source could have the second amplitude and opposite phase such that when current source 202 "pushes" current into patient 208 through electrode 206a, the second current source "pulls" current from the patient through electrode 206b. As will be apparent to those skilled in the art, this can provide a virtual ground in the patient.

In various other embodiments, a greater number of electrodes could be used for current injection/return and voltage measurement. For example, but not limited to, a four point measurement could be used. A four point measurement can use four electrodes, where one electrode is used as a current injection electrode, a second electrode could be used as a current return electrode, a third electrode could be used as a positive voltage measurement electrode, and a fourth electrode could be used as a negative voltage measurement electrode.

Referring back to FIG. 3, electrode 206a could be used as used as voltage measurement electrode. Thus, it will be appreciated that in this example, the current delivery circuit **220***b* is actually acting to deliver a sensed voltage signal. It will be appreciated from this that the circuit can be more generally referred to as a signal delivery circuit. In this example, the apparatus of FIG. 3 can be combined with a current return electrode and a second voltage measurement electrode so a four point measurement system would result, for example to allow tetrapolar impedance measurements to be performed.

In various embodiments, an AC current such as sinusoidal waveform with no DC offset can be used and therefore in such embodiments the application of the terms current injection and current return to an electrode is arbitrary. More specifically, in such an embodiment, a given electrode will be conducting current in one direction half of the time and conducting it in the other direction for the other half. For similar reasons; the application of the terms positive and negative voltage measurement to an electrode can also be arbitrary.

In addition, when a four point measurement is used, a more accurate voltage measurement may be obtained. Specifically, substantially no current will be moving through the voltage measurement electrodes and therefore the impedance of the electrodes and the interface between the electrode and patient will not significantly distort the voltage measurement.

Reference is now made to FIG. 4, which a block diagram illustrating a portion of the current multiplexing system 200 of FIG. 2 in greater detail. Multiplexer 204 causes a large parasitic impedance to appear at its outputs such as at node 222. Electrode 206 is coupled to the current delivery circuit at node 224. Node 226, which is the output of current delivery circuit 220 can be used to measure the voltage that appears at electrode 206. Optionally, node 226 can be connected to the input of a multiplexer, which may be useful when there are many electrodes 206 and current delivery circuits 220 being used.

In some embodiments, current delivery circuit 220 comprises a current buffer 430 and a voltage buffer 440. Current buffer 430 introduces parasitic impedance 438 to appear at node 224. In some embodiments, current buffer 430 can be a current conveyor as shown in FIG. 4. The input 432 of current buffer 430 is coupled to node 222. The output 434 of current buffer 430 is coupled to node 224. If a current conveyor is used as the current buffer then its other input 436 can be connected to ground.

In various embodiments according to applicants' teachings, current buffer 430 is designed so as to have a low input impedance at input 432. Consequently, the leakage current (ΔI_1) lost to parasitic impedance **410** is minimal. In addition, current buffer 430 is designed so as to present only a small 5 parasitic impedance to node 224. As a result most of the current that is outputted through output 434 passes through electrode 206 and only a minimal amount of leakage current (ΔI_2) is lost to parasitic impedance 438. In addition, in various embodiments, current buffer 430 is designed to have 10 a high output impedance, which maximizes the amount of current delivered to the patient electrode. The combination of the above features can be achieved for example by implementing current buffer 430 as a current conveyor. However, in some other embodiments other appropriate 15 current buffers with similar, features can also be used.

In some embodiments, voltage buffer **440** can be selected to have a high input impedance so that the voltage at node **224** can be reproduced at node **226** without affecting the voltage at node **224**. In various embodiments, this allows for 20 standardized non-invasive voltage measurements to be made at node **226**.

In various embodiments, current multiplexing system **200** can be used without a calibration step even when the loads are changed. For example, a calibration step is not required 25 when a electrodes **206** is coupled to a new, patient. In addition, in various embodiments, current multiplexing system **200** can be used without modifying or otherwise manipulating the patient electrode interface.

In various embodiments, current delivery circuit **220** can ³⁰ be integrated on a single integrated circuit. In some embodiments, a plurality of current delivery circuit **220** can be integrated on a single integrated circuit and packaged in a single chip.

It should be understood that in various embodiments any 35 appropriate scheme could be used to switch or multiplex various inputs to each current injection circuit **220**. Regardless of the switching or multiplexing scheme utilized a resulting parasitic capacitance **410** will result. Current buffer **430** isolates the electrode patient interface from interfering 40 effects caused by the switches and their parasitic elements. Similarly, voltage buffer **440** isolates electrode **206** (and thereby the patient electrode interface) from interfering effects that may otherwise be caused by components beyond the voltage buffer **440**. 45

Accordingly, in one example, the apparatus can be used to supply a drive signal to an electrode attached to the biological subject, and sense a signal at the electrode. Whilst the above example has focussed on the delivery of a current and measurement of a voltage, this is not essential, and the drive 50 signal could be a voltage signal, with a current signal being measured. Alternatively, the signal delivery circuit could be used to measure the drive signal applied to the subject, allowing this to be used for example in calculating impedance measurements or the like. This may be performed to 55 ensure that the magnitude and phase of the drive signal which is used in impedance calculations is as accurate as possible.

Accordingly, in general terms, the apparatus includes a signal delivery circuit including first and second buffers. The 60 first buffer includes a first buffer input for receiving a signal from a signal source and a first buffer output for supplying a drive signal to an electrode attached to the biological subject. Similarly, the second buffer includes a second buffer input coupled to the first buffer output and a second buffer 65 output for providing a sensed signal indicative of a signal at the electrode, to a sensor.

In one example, the first buffer is a current buffer and the second buffer a voltage buffer, allowing a drive current to be applied to the subject, and to allow a voltage at the electrode to be sensed. However, in alternative examples, a voltage signal may be applied to the subject and a current flow through the subject measured.

The signal delivery circuit allows a drive signal to be applied to the subject via an electrode and then a signal measured at the electrode. In one example, the measured signal is indicative of a signal, such as a potential, across the biological subject, with a single electrode is used as both a drive a sense electrode. However, this is not essential, and in another example, first and second current delivery circuits can be coupled to drive and sense electrodes respectively.

Accordingly, in one example, the first signal delivery circuit can include a first buffer having a first buffer input for receiving a signal from the signal source and a first buffer output for supplying the drive signal to the drive electrode and a second buffer having a second buffer input coupled to the first buffer output and a second buffer output for providing a sensed signal indicative of the drive signal at the drive electrode. This allows the magnitude of the drive signal injected into the biological subject to be measured, allowing this value to be used in performing impedance determination or the like. In this example, if the drive signal is a current signal, the second buffer is typically an amplifier having inputs in parallel with a resistor positioned between the first buffer output and the electrode, thereby allowing the current injected into the subject to be measured.

In this example, the second signal delivery circuit can include a first buffer having a first buffer input coupled to ground and a first buffer output coupled to the sense electrode and a second buffer having a second buffer input coupled to the first buffer output and a second buffer output for providing a sensed signal indicative of a signal at the sense electrode.

Reference is now made to FIG. 5, which is a schematic diagram of the current multiplexing system of FIG. 4 showing additional components. In various embodiments, current delivery circuit 200 can also comprise a DC offset control circuit 550. In some embodiments, DC offset control circuit 550 is coupled in a feedback loop of the current buffer 430. More specifically, DC offset control circuit 550 is connected between node 226 and node 222. In various embodiments, DC offset or DC bias at node 224. However, DC offset control circuit 550 is coupled to node 222 so as not to disturb the electrode-patient interface which exists between electrode 206 and patient 208. This use of DC offset control circuit 550 can increase the accuracy of the measurements made using current multiplexing system 200.

In various embodiments, current delivery circuit **200** can also comprise a negative impedance circuit **570**. Negative impedance circuit **570** can be used to compensate for parasitic impedance **438** introduced by current buffer **430** or other circuit elements at, for example, node **224**. Negative impedance circuit **570** comprises an amplifier **571** and a compensation impedance **591**. In various embodiments, voltage amplifier **571** has a gain value greater than 1. Negative impedance circuit **570** can be used to provide a current to node **224** that is substantially equal to the leakage current (ΔI_2) that is lost from node **224** through parasitic impedance **438**. The operation of negative impedance circuit **570** will be discussed in greater detail below.

It should be understood that the use of DC offset control circuit **550** is optional. It should also be understood that the use of negative impedance circuit **570** is optional. It should

be further understood that DC offset control circuit 550 and negative impedance circuit 570 can be used independently of each other. Therefore, various embodiments can utilize either of one DC offset control circuit 550 and negative impedance circuit 570, both DC offset control circuit 550⁻⁵ and negative impedance circuit 570, or neither.

It should further be understood that in various embodiments, DC offset control circuit 550 and negative impedance circuit 570 can be integrated as part of current delivery circuit 220. In various other embodiments, either or both of DC offset control circuit 550 and negative impedance circuit 570 can be separate circuits. For example, but not limited to, in some embodiments, multiplexers could be used to couple either of these circuit to a particular current delivery circuit. 15

Reference is now made to FIG. 6, which is a schematic diagram of a circuit 600 comprising a transmission channel 601, a parasitic impedance 438 shunting the transmission line, and a negative impedance circuit 570. Transmission channel 601 can, for example, correspond to node 224. 20 Negative impedance circuit 570 comprises a voltage amplifier 571 and compensation impedance 591. The input of the amplifier 571 is coupled to the signal-transmission channel 601 and the output is coupled to one terminal of the compensation impedance 591. The other terminal of the 25 compensation impedance is coupled to the signal-transmission channel.

Signal-transmission channel 601 may be used to transmit a current to a load (not shown), which may be any suitable circuit or circuit component, such as for example an electrode. The presence of a signal on signal-transmission channel 601 causes a voltage to appear across parasitic impedance 438. This causes a leakage current $I_{leakage}$ to flow through the parasitic impedance 438. The magnitude of the 35 current flowing through parasitic impedance 438 depends on the value of the impedance as well as the magnitude of the voltage appearing across its terminals.

Amplifier 571 amplifies the signal appearing on the signal-transmission channel 601. In various embodiments, 40 amplifier 571 has a gain that is greater than 1. This causes a voltage to appear across compensation impedance 591 and a current I_{comp} to flow through compensation impedance **591**.

In various embodiments, the gain of amplifier 571 and the $_{45}$ value of the compensation impedance is selected such that the current that flows through parasitic impedance 438 is compensated for by the current that flows through compensation impedance **591**. Specifically, given a signal voltage of V_{signal} , a parasitic impedance of Z_{para} , the leakage current 50 can be said to be:

$$I_{leakage} = V_{signal} \times \left(\frac{1}{Z_{para}}\right)$$
 Equation (1)

Similarly, given a compensation impedance of Z_{comp} and an amplifier gain of G, the compensation current flowing through the compensation current may be said to be:

$$I_{comp} = V_{signal} \times (G-1) \times \left(\frac{1}{Z_{comp}}\right)$$
 Equation (2)

65

Equating equation (1) and equation (2) yields the following:

 $I_{comp} = I_{leakage}$

 $\left(\frac{G-1}{Z_{comp}}\right) = \frac{1}{Z_{para}}$

Equation (3)

Thus, by selecting G and Z_{comp} to satisfy equation (3) the compensation current will exactly match the leakage current. The compensation impedance 591 effectively serves as a negative impedance that cancels the effect of the parasitic impedance 438.

In various embodiments, the value of the parasitic impedance may not be known and therefore it may not be possible to select a gain for the amplifier by simply using equation (3)above. In such embodiments, the value of the gain can be estimated by using circuit 600 of FIG. 6. Specifically, circuit 600 is implemented by selecting a compensation impedance and range of values of gain. The circuit is operated at the various values of gain and the output is monitored. For those values of gain that exceed the required value the output would oscillate. Thus, the correct value of the gain lies in a range of values that is bounded by (1) the lowest known value of the gain at which the output oscillates and (2) the highest known value of the gain at which the output does not oscillate. This process may be continued in an iterative manner until a suitable value of gain is selected. Once an appropriate value of gain is determined, the parasitic impedance may be estimated by using equation (3) given above.

In various embodiments, the parasitic impedance may be comprised of both capacitive and resistive elements. However, in some embodiments the effect of the capacitive loading can be significantly greater than the effect of the resistive loading. In such cases, various embodiments of applicants' teachings may be used to address the capacitive loading and not the resistive loading. Alternatively, applicants' teachings may be used to partially compensate for any portion of the parasitic impedance. Thus, in various embodiments, circuits according to applicants' teachings may be used to reduce and partially compensate for any leakage currents that may flow through any parasitic impedances coupled to a signal-transmission channel, but not necessarily to completely compensate for all the current that is lost due to leakage currents.

Alternatively, the parasitic impedance may be measured or estimated according to known techniques. The value of the parasitic impendence obtained from this may then be used to select initial values for the compensation impedance and the range of values of gain. The gain can then be fine tuned according to the above-described method.

Reference is now made to FIG. 7, which is a schematic diagram of the current multiplexing system of FIG. 5, showing the DC bias control and negative impedance in greater detail. In various embodiments, voltage buffer 440 55 comprises an operational amplifier 742 connected as an output follower. More specifically, non-inverting input 744 is coupled to node 224; while, inverting node 746 and output 748 are coupled to node 226.

In various embodiments, negative impedance circuit 570 60 comprises an operational amplifier 772 with non-inverting input 774, inverting input 776 and output 778. Resistor 780 is connected between node 226 and non-inverting input 774. Resistor 782 is coupled between inverting input 776 and ground. One terminal of resistor 784 is coupled to output 778 and the other terminal of resistor 784 is coupled to resistor 786. Capacitor 788 and resistor 790 are each coupled between non-inverting input 776 and the common terminal

of resistors 784 and 786. Capacitor 792 and resistor 794 are coupled between node 224 and resistor 786. Negative impedance circuit will be discussed in greater detail below.

In some embodiments DC-offset control circuit 550 comprises an operational amplifier 752 connected as an integra- 5 tor. Specifically, operational amplifier 752 has non-inverting input 754, inverting input 756 and output 758. Capacitor 760 is connected between inverting input 756 and output 758. Resistor 762 is connected between inverting input 756 and node 226. Resistor 764 is connected between the non- 10 inverting input 754 and ground. Resistor 766 is connected between output 758 and node 222.

Reference is now made to FIG. 8, which is a schematic diagram of negative impedance circuit 570 of FIG. 7. For greater clarity FIG. 8 illustrates negative impedance circuit 15 570 apart from the rest of current multiplexing system 200.

Negative impedance circuit 570 comprises an amplifier portion 571 having an operational amplifier 772 with a non-inverting input 774, an inverting input 776, an output node 778. Operational amplifier 772 also comprises power 20 rails 895 and 896.

Referring again to the amplifier portion 571, amplifier portion 571 further comprises an input balancing portion 897, a gain control portion 898, and a stability control portion 899. Input balancing portion 897 comprises resistor 25 780, which is connected between node 226 and non-inverting input 774. Gain control portion 898 comprises resistor 782 and resistor 790. By adjusting the values of resistors 782 and 790, one is able to adjust the gain G of the overall amplifier portion 771. In various embodiments, the values of 30 resistors 782 and 790 may be set to a value that is greater than 1. Stability control portion 899 comprises capacitor 788, resistor 784, and resistor 794. By adjusting the values of capacitor 788, resistor 784, and resistor 794 one is able to alter the stability of the overall amplifier circuit.

In various embodiments, negative impedance circuit 570 also comprises compensation impedance 591. Compensation impedance 591 is in turn comprised of resistor 794 and capacitor 792, both of which are connected between node 224 and stability control portion 899. Compensation imped- 40 ance 591 is used to compensate for parasitic impedance 438 of FIG. 7. By adjusting gain control portion 898 and compensation impedance 591, one may adjust the compensation current that is provided to the signal-transmission channel, and thereby match the compensation current mag- 45 nitude to the magnitude of the leakage current. This may be done according to equation (3) given above.

Whilst the above examples have focussed on the use of a current source, this is not essential, and alternatively the system may use a voltage source, with the generated voltage 50 being used to inject a drive signal in the form of a current into a subject, with the magnitude of the drive signal being measured using a sensor at node 214.

From this, it will be appreciated that the current delivery circuit 220 can be referred to more generally as a signal 55 delivery circuit. Similarly, the current and voltage buffers can be first and second buffers for supplying a drive signal to an electrode attached to the biological subject, and for providing a signal indicative of a signal at the electrode.

As mentioned above, the signal delivery circuit can be 60 used in a variety of different systems. In one example, the signal delivery circuit can be incorporated into apparatus suitable for performing an analysis of a subject's bioelectric impedance, an example of which will now be described with reference to FIG. 9.

As shown the apparatus includes a measuring device 900 including a processing system 902, connected to one or more

65

signal generators 917A, 917B, via respective first leads 923A, 923B, and to one or more sensors 918A, 918B, via respective second leads 925A, 925B.

In use, the signal generators 917A, 917B are coupled to two first electrodes 913A, 913B, via drive signal delivery circuits 919A, 919B, which therefore act as drive electrodes to allow signals to be applied to the subject S, whilst the one or more sensors 918A, 918B are coupled to the second electrodes 915A, 915B, via sensed signal delivery circuits 920A, 920B, which act as sense electrodes, allowing signals across the subject S to be sensed. It will be appreciated that the drive and sensed signal delivery circuits are similar to the delivery circuits described above in FIGS. 1 to 8.

In one example, a single signal generator 917 may be provided, coupled to the drive signal delivery circuits 919A, 919B and hence the drive electrodes 913A, 913B, via multiplexers 204a from the example of FIG. 3. Similarly a single sensor 918 can be coupled to the sensed signal delivery circuits 920A, 920B and hence the sense electrodes **915**A, **915**B, via multiplexers, such as the multiplexers **212**b from the example of FIG. 3.

However, this is not essential, and alternatively first and second signal generators 917A, 917B and sensors 918A, 918B can be used, each being coupled to the corresponding electrodes 913A, 913B, 914A, 914B via respective signal delivery circuits. This is particularly useful if the arrangement is used to perform balancing, as will be described in more detail below.

Accordingly, this provides apparatus for use in performing impedance measurements on a subject which includes a signal source for applying a signal to a subject, a sensor for sensing signals from the subject, a processing system for controlling the signal source and receiving signals from the sensor and at least one signal delivery circuit, including a first buffer having: a first buffer input for receiving a signal from a signal source and a first buffer output for supplying a drive signal to an electrode attached to the biological subject and a second buffer having a second buffer input coupled to the first buffer output and a second buffer output for providing a sensed signal indicative of a signal at the electrode, to a sensor.

In one example, at least one signal delivery circuit is used for supplying a current to a drive electrode attached to the biological subject and at least one signal delivery circuit for providing a voltage signal indicative of a voltage at a sense electrode.

Additional features of the impedance measurement apparatus will now be described.

The signal generators 917A, 917B and the sensors 918A, 918B may be provided at any position between the processing system 902 and the electrodes 913A, 913B, 915A, 915B, and may be integrated into the measuring device 900. However, in one example, the signal generators 917A, 917B and the sensors 918A, 918B are integrated into an electrode system, or another unit provided near the subject S, with the leads 923A, 923B, 925A, 925B connecting the signal generators 917A, 917B and the sensors 918A, 918B to the processing system 902.

It will be appreciated that the above described system is a two channel device, used to perform a classical fourterminal impedance measurement, with, each channel being designated by the suffixes A, B respectively. The use of a two channel device is for the purpose of example only, as will be described in more detail below.

An optional external interface 903 can be used to couple the measuring device 900, via wired, wireless or network connections, to one or more peripheral devices 904, such as an external database or computer system, barcode scanner, or the like. The processing system 902 will also typically include an I/O device 905, which may be of any suitable form such as a touch screen, a keypad and display, or the like.

In use, the processing system 902 is adapted to generate control signals, which cause the signal generators 917A, 917B to generate one or more alternating signals, such as voltage or current signals of an appropriate waveform, which can be applied to a subject S, via the first electrodes 10 913A, 913B. The sensors 918A, 918B then determine the voltage across or current through the subject S, using the second electrodes 915A, 915B and transfer appropriate signals to the processing system 902, for analysis. In the event that the apparatus includes multiplexers for coupling 15 signal generators or sensors to respective signal delivery circuits, the processing system would also typically act to control the multiplexers allowing signals to be delivered to and measured across the subject as required.

Accordingly, it will be appreciated that the processing 20 system 902 may be any form of processing system which is suitable for generating appropriate control signals and at least partially interpreting the measured signals to thereby determine the subject's bioelectrical impedance, and optionally determine other information such as relative fluid levels, 25 impedance of the system at the applied frequencies. One or the presence, absence or degree of conditions, such as oedema, lymphoedema, measures of body composition, cardiac function, or the like.

The processing system 902 may therefore be a suitably programmed computer system, such as a laptop, desktop, 30 PDA, smart phone or the like. Alternatively the processing system 902 may be formed from specialised hardware, such as an FPGA (field programmable gate array), or a combination of a programmed computer system and specialised hardware, or the like, as will be described in more detail 35 helow

In use, the first electrodes 913A, 913B are positioned on the subject to allow one or more signals to be injected into the subject S. The location of the first electrodes will depend on the segment of the subject S under study. Thus, for 40 example, the first electrodes 913A, 913B can be placed on the thoracic and neck region of the subject S to allow the impedance of the chest cavity to be determined for use in cardiac function analysis. Alternatively, positioning electrodes on the wrist and ankles of a subject allows the 45 impedance of limbs and/or the entire body to be determined, for use in oedema analysis, or the like.

Once the electrodes are positioned, one or more alternating signals are applied to the subject S, via the first leads 923A, 923B and the first electrodes 913A, 913B. The nature 50 of the alternating signal will vary depending on the nature of the measuring device and the subsequent analysis being performed.

For example, the system can use Bioimpedance Analysis (BIA) in which a single low frequency signal (typically <50 55 kHz) is injected into the subject S, with the measured impedance being used directly in the assessment of relative intracellular and extracellular fluid levels. In contrast Bioimpedance Spectroscopy (BIS) devices utilise frequencies ranging from very low frequencies (4 kHz) to higher 60 frequencies (1000 kHz), and can use as many as 256 or more different frequencies within this range, to allow multiple impedance measurements to be made within this range.

Thus, the measuring device 900 may either apply an alternating signal at a single frequency, at a plurality of 65 frequencies simultaneously, or a number of alternating signals at different frequencies sequentially, depending on the

preferred implementation. The frequency or frequency range of the applied signals may also depend on the analysis being performed.

In one example, the applied signal is generated by a voltage generator, which applies an alternating voltage to the subject S, although alternatively current signals may be applied. In one example, the voltage source is typically symmetrically arranged, with each of the signal generators 917A, 917B being independently controllable, to allow the signal voltage across the subject to be varied.

A voltage difference and/or current is measured between the second electrodes 915A, 915B. In one example, the voltage is measured differentially, meaning that each sensor 918A, 918B is used to measure the voltage at each second electrode 915A, 915B and therefore need only measure half of the voltage as compared to a single ended system.

The acquired signal and the measured signal will be a superposition of voltages generated by the human body, such as the ECG (electrocardiogram), voltages generated by the applied signal, and other signals caused by environmental electromagnetic interference. Accordingly, filtering or other suitable analysis may be employed to remove unwanted components.

The acquired signal is typically demodulated to obtain the suitable method for demodulation of superposed frequencies is to use a Fast Fourier Transform (FFT) algorithm to transform the time domain data to the frequency domain. This is typically used when the applied current signal is a superposition of applied frequencies. Another technique not requiring windowing of the measured signal is a sliding window FFT.

In the event that the applied current signals are formed from a sweep of different frequencies, then it is more typical to use a signal processing technique such as multiplying the measured signal with a reference sine wave and cosine wave derived from the signal generator, or with measured sine and cosine waves, and integrating over a whole number of cycles. This process, known variously as quadrature demodulation or synchronous detection, rejects all uncorrelated or asynchronous signals and significantly reduces random noise.

Other suitable digital and analogue demodulation techniques will be known to persons skilled in the field.

In the case of BIS, impedance or admittance measurements are determined from the signals at each frequency by comparing the recorded voltage and the current through the subject. The demodulation algorithm can then produce amplitude and phase signals at each frequency.

As part of the above described process, the distance between the second electrodes 915A, 915B may be measured and recorded. Similarly, other parameters relating to the subject may be recorded, such as the height, weight, age, sex, health status, any interventions and the date and time on which they occurred. Other information, such as current medication, may also be recorded. This can then be used in performing further analysis of the impedance measurements, so as to allow determination of the presence, absence or degree of oedema, to assess body composition, or the like.

The accuracy of the measurement of impedance can be subject to a number of external factors. These can include, for example, the effect of capacitive coupling between the subject and the surrounding environment, the leads and the subject, the electrodes, or the like, which will vary based on factors such as lead construction, lead configuration, subject position, or the like. Additionally, there are typically variations in the impedance of the electrical connection between

the electrode surface and the skin (known as the "electrode impedance"), which can depend on factors such as skin moisture levels, melatonin levels, or the like. A further source of error is the presence of inductive coupling between different electrical conductors within the leads, or between 5 the leads themselves.

Such external factors can lead to inaccuracies in the measurement process and subsequent analysis and accordingly, it is desirable to be able to reduce the impact of external factors on the measurement process.

One form of inaccuracy that can arise is caused by the voltages across the subject being unsymmetrical, a situation referred to as an "imbalance". Such a situation results in a significant signal voltage at the subject's body centre, which in turn results in stray currents arising from parasitic capaci-15 tances between the subject's torso and the support surface on which the subject is provided.

The presence of an imbalance, where the voltage across the subject is not symmetrical with respect to the effective centre of the subject, leads to a "common mode" signal, 20 which is effectively a measure of the signal at the subject S that is unrelated to the subject's impedance.

To help reduce this effect, it is therefore desirable for signals to be applied to the subject S that they result in a symmetrical voltage about the subject's body centre. As a 25 result, a reference voltage within the subject S, which is equal to a reference voltage of the measurement apparatus, will be close to the effective body centre of the subject, as considered relative to the electrode placement. As the measuring device reference voltage is typically ground, this 30 results in the body centre of the subject S being as close to ground as possible, which minimises the overall signal magnitude across the subject's torso, thereby minimising stray currents.

In one example, a symmetrical voltage about the sensing 35 electrodes can be achieved by using a symmetrical voltage source, such as a differential bidirectional voltage drive scheme, which applies a symmetrical voltage to each of the drive electrodes **913A**, **913**B. However, this is not always effective if the contact impedances for the two drive elec- 40 trodes **913A**, **913**B are unmatched, or if the impedance of the subject S varies along the length of the subject S, which is typical in a practical environment.

In one example, the apparatus overcomes this by adjusting the differential voltage drive signals applied to each of 45 the drive electrodes **913**A, **913**B, to compensate for the different electrode impedances, and thereby restore the desired symmetry of the voltages across the subject S. This process is referred to herein as balancing and in one example, helps reduce the magnitude of the common mode 50 signal, and hence reduce current losses caused by parasitic capacitances associated with the subject.

The degree of imbalance, and hence the amount of balancing required, can be determined by monitoring the signals at the sense electrodes **915**A, **915**B, and then using 55 these signals to control the signal applied to the subject via the drive electrodes **913**A, **913**B. In particular, the degree of imbalance can be calculated by determining an additive voltage from the voltages detected at the sense electrodes **915**A, **915**B. 60

In one example process, the voltages sensed at each of the sense electrodes **915**A, **915**B are used to calculate a first voltage, which is achieved by combining or adding the measured voltages. Thus, the first voltage can be an additive voltage (commonly referred to as a common mode voltage 65 or signal) which can be determined using a differential amplifier.

In this regard, a differential amplifier is typically used to combine two sensed voltage signals V_a , V_b , to determine a second voltage, which in one example is a voltage differential V_a-V_b across the points of interest on the subject S. The voltage differential is used in conjunction with a measurement of the current flow through the subject to derive impedance values. However, differential amplifiers typically also provide a "common mode" signal $(V_a+V_b)/2$, which is a measure of the common mode signal.

Whilst differential amplifiers include a common mode rejection capability, this is generally of only finite effect and typically reduces in effectiveness at higher frequencies, so a large common mode signal will produce an error signal superimposed on the differential signal.

The error caused by common mode signals can be minimised by calibration of each sensing channel. In the ideal case where both inputs of a differential amplifier are perfectly matched in gain and phase characteristics and behave linearly with signal amplitude, the common mode error will be zero. In one example, the two sensing channels of the differential amplifier are digitised before differential processing. It is therefore straightforward to apply calibration factors independently to each channel to allow the characteristics to be matched to a high degree of accuracy, thereby achieving a low common mode error.

Accordingly, by determining the common mode signal, the applied voltage signals can be adjusted, for example by adjusting the relative magnitude and/or phase of the applied signals, to thereby minimise the common mode signal and substantially eliminate any imbalance.

An example of the operation of the apparatus of FIG. **9** to perform this will now be described with reference to FIG. **10**.

At step 1000, a first signal is applied to the subject S, with a second signal measured across the subject S being determined at step 1010. This will typically be achieved using the techniques outlined above. Accordingly, the processing system 902 will cause the signal generators 917A, 917B to generate the first signal, which is typically applied to the subject S via the first electrodes 913A, 913B. Similarly the second signal will be sensed by the sensors 918A, 918B, via the second electrodes 915A, 915B, with an indication of the second signal being provided to the processing system 902.

At step **1020**, an imbalance is determined by the processing system **902** using the second signal sensed at the second electrodes **915**A, **915**B, which in one example represents a common mode signal.

At step **1030**, the measuring device optionally adjusts the first signal applied to the subject S, so as to reduce the imbalance and hence the magnitude of the common mode signal. Thus, the magnitude of the signal applied at either one of the first electrodes **913A**, **913B** can be adjusted, for example by increasing or decreasing the relative signal magnitudes and/or altering the relative signal phases, so as to balance the signal within the subject and centralise the position of the reference voltage within the subject relative to the electrode positioning.

At step **1040**, the measuring device can then determine the signal applied to the subject and the voltages measured at the electrodes **913A**, **913B**, thereby allowing an impedance to be determined at step **1050**.

As the position of the reference voltage within the subject S is impedance dependent, the imbalance will typically vary depending on the frequency of the applied signal. Accordingly, in one example, it is typical to determine the imbalance and adjust the applied signal at each applied frequency. However, this may depend on the preferred implementation.

An example of balancing procedure and operation of an impedance measuring device is described in more detail in the copending patent application number WO2009/059351, and this will not therefore be described in any further detail herein.

In various examples, applicants' teachings relate to a current delivery circuit. In various examples, the current delivery circuit comprises an input, a current output and a voltage output. In some examples, the current delivery circuit accepts a current at its input and produces a proportional output current at the current output. In various examples, current delivery circuit produces an output voltage signal at the voltage output that is proportional to an input voltage signal appearing at the current delivery circuit current output.

In various examples according to applicants' teachings, the current delivery circuit comprises a current buffer and a voltage buffer. The current buffer and voltage buffer each have an input and an output. The current delivery circuit 20 input is coupled to the current buffer input. The current buffer output is coupled to the current delivery circuit current output. The voltage buffer input is coupled to the current delivery circuit current output. The voltage buffer output is coupled to the current delivery circuit voltage 25 output.

In some examples, the output current is substantially equal to the input current. In various examples, the output voltage signal is substantially equal to the input voltage signal.

In some examples, the current buffer is a current conveyor. In various examples, the voltage buffer is a voltage follower.

In various examples, the current delivery circuit further comprises a parasitic impedance coupled to the current 35 buffer output. In some examples, the current delivery circuit further comprises a negative impedance coupled between, the voltage buffer output and the current buffer output for compensating for the parasitic impedance. In some examples, the negative impedance comprises: a compensa- 40 tion impedance having a compensation impedance value, a first terminal and a second terminal, the first terminal coupled to the voltage buffer input; a compensation amplifier having a compensation amplifier input, a compensation amplifier output and a gain, the compensation amplifier 45 input coupled to the voltage buffer output, the compensation amplifier output coupled to the second terminal of the compensation impedance to provide a compensation current to flow through the impedance, and the gain and the compensation impedance values are selected based on the para- 50 sitic impedance value so that the compensation current has a magnitude substantially equal to the leakage current magnitude.

In various examples, current delivery circuit further comprises a DC offset control circuit coupled between the 55 age output is coupled a first voltage multiplexer and the voltage buffer output and the current buffer input. In some examples according to applicants' teachings, the DC offset control comprises an integrator coupled between the voltage buffer output and the current buffer input.

In various examples, applicants' teachings relate to a 60 current multiplexing circuit comprising one or more current multiplexers, each having an input and a plurality of outputs, and one or more current delivery circuits. In various examples, the outputs of the one or more current multiplexers are coupled to the input of each of the one or more 65 current delivery circuits. In some examples, the current multiplexing circuit further comprises one or more voltage

multiplexers. The voltage output of each of the one or more current delivery circuits is coupled to an input of the one or more voltage multiplexers.

In various examples, the circuit further comprises one or more current sources. In various examples, each current source is coupled to the input of the one or more current multiplexers.

In various examples, applicants' teachings relate to a current multiplexing system. In various examples, current multiplexing system comprises a first current delivery circuit and a second current delivery circuit. Current multiplexing system further comprises a first electrode coupled to the first current delivery circuit current output and a second electrode coupled to the second current delivery circuit current output.

In various examples, the voltage output of the first current delivery circuit is coupled to an input of a first voltage multiplexer and the voltage output of the second current delivery circuit is coupled to an input of a second voltage multiplexer. In various examples, the outputs of the first and second voltage multiplexers are adapted to provide first and second voltage measurement signals for allowing a differential voltage measurement.

In some examples, a first current source is coupled to the input of the first current delivery circuit. In some examples, the first current source is controllable to provide a constant alternating current. In some examples, the constant alternating current has a constant amplitude and constant phase. In various examples, the amplitude and phase are controllably variable.

In some examples, the input of the second current delivery circuit is coupled to ground. In various other examples, the input of the second current delivery circuit is coupled to a second current source. In various examples the second current source is complimentary to the first current source. As used herein, the term complimentary current source refers to a current source providing a current of equal magnitude with a 180° phase shift.

In some examples, the current multiplexing system further comprises one or more current multiplexers. Each current multiplexer has a first output coupled to the first current delivery circuit and a second output coupled to the second current delivery circuit. In some examples, the input of at least one of the current multiplexers is coupled to a first current source. In some examples, the input of at least one of the current multiplexers is coupled to a second current source. In some examples, the input of at least one of the current multiplexers is coupled to ground.

In some examples, the current multiplexing system further comprises a third and fourth current delivery circuit as well as a third and fourth electrode. The third electrode is coupled to the third current delivery circuit current output. The fourth electrode is coupled to the fourth current delivery circuit current output.

In some examples, the third current delivery circuit voltfourth current delivery circuit voltage output is coupled a second voltage multiplexer. In various examples, the outputs of the first and second voltage multiplexers are adapted to provide first and second voltage measurement signals for allowing a differential voltage measurement.

In some examples, the third and forth current delivery circuit inputs are coupled to ground.

In some examples of applicants' teachings, current multiplexing system comprises a current source, a current multiplexer, a current delivery circuit, an electrode, and an voltage multiplexer. The current source is coupled to an input of the current multiplexer. An output of the current

multiplexer is coupled to the input of the current delivery circuits. A current output of the current delivery circuit is coupled to the electrode. A voltage output of the current delivery circuit is coupled to an input of the voltage multiplexer. In various examples the current multiplexing system 5 further comprises an electrode coupled to the current output of the current delivery circuit.

Persons skilled in the art will appreciate that numerous variations and modifications will become apparent. All such variations and modifications which become apparent to 10 persons skilled in the art, should be considered to fall within the spirit and scope that the invention broadly appearing before described.

Features from different examples above may be used interchangeably or in conjunction, where appropriate. Thus, 15 for example, a range of different techniques are described for minimising errors and these can be used independently of each other, or in conjunction, depending on the particular implementation.

Furthermore, whilst the above examples have focussed on 20 a biological subject such as a human, it will be appreciated that the measuring device and techniques described above can be used with any animal, including but not limited to, primates, livestock, performance animals, such as race horses, or the like. 25

The above described processes can be used for diagnosing the presence, absence or degree of a range of conditions and illnesses, including, but not limited to oedema, lymphoedema, body composition, or the like.

The claims defining the invention are as follows:

1. Apparatus for electrically connecting measurement apparatus to a biological subject, the apparatus including a signal delivery circuit including: 35

a current buffer having:

- a current buffer input for receiving a signal from a signal source; and,
- a current buffer output for supplying a current to an electrode attached to the biological subject;

a voltage buffer having:

- a voltage buffer input coupled to the current buffer output; and,
- a voltage buffer output for providing a voltage signal indicative of a voltage at the electrode, to a sensor; and
- a DC offset control circuit coupled between the voltage buffer output and the current buffer input.

2. Apparatus according to claim 1, wherein the current buffer is a current conveyor.

3. Apparatus according to claim 1, wherein the voltage 50 signal source is coupled to ground. buffer is an amplifier connected as an output follower.

4. Apparatus according to claim 1, wherein the DC offset control circuit includes an integrator coupled between the voltage buffer output and the current buffer input.

control circuit is used to control a DC offset at the electrode.

6. Apparatus according to claim 1, wherein the signal delivery circuit includes a negative impedance circuit coupled between the voltage buffer output and the current buffer output. 60

7. Apparatus according to claim 6, wherein the negative impedance circuit includes a negative impedance circuit amplifier and a compensation impedance.

8. Apparatus according to claim 7, wherein at least one of a gain of the negative impedance circuit amplifier and a 65 value of the compensation impedance is selected to compensate for parasitic impedance losses.

9. Apparatus according to claim 6, wherein the negative impedance circuit includes:

- a compensation impedance having a compensation impedance value, a first terminal and a second terminal, the first terminal coupled to the current buffer output; and.
- a compensation amplifier having a compensation amplifier input, a compensation amplifier output and a gain, the compensation amplifier input coupled to the voltage buffer output, the compensation amplifier output coupled to the second terminal of the compensation impedance to provide a compensation current to flow through the impedance, and the gain and the compensation impedance values are selected based on a parasitic impedance value so that the compensation current has a magnitude substantially equal to a leakage current magnitude.

10. Apparatus according to claim 1, wherein the apparatus includes

- a plurality of signal delivery circuits, each signal delivery circuit being for supplying a current to a respective electrode attached to the biological subject;
- at least one signal source; and,
- at least one first multiplexer for selectively connecting the signal source to one of the plurality of signal delivery circuits to thereby apply a current to the biological circuit via the respective electrode.

11. Apparatus according claim 1, wherein the apparatus includes:

a plurality of signal delivery circuits, each signal delivery circuit being for providing a voltage signal indicative of a voltage at the electrode, to a sensor;

at least one sensor; and,

at least one multiplexer for selectively connecting the at least one sensor to one of the plurality of signal delivery circuits to thereby allow the sensor to sense the voltage signal indicative of a voltage at the respective electrode.

12. Apparatus according to claim 1, wherein the apparatus 40 includes, first and second signal sources for generating respective first and second drive signals, the first and second signal sources being coupled to respective signal delivery circuits.

13. Apparatus according to claim 12, wherein the first and 45 second signal sources are for generating at least one of:

complementary signals;

signals having a constant amplitude and phase; and,

signals having a controlled amplitude and phase.

14. Apparatus according to claim 1, wherein at least one

15. Apparatus according to claim 1, wherein the measurement apparatus includes at least one signal source and at least one sensor.

16. Apparatus according to claim 15, wherein the mea-5. Apparatus according to claim 1, wherein the DC offset 55 surement apparatus is an impedance measurement apparatus.

17. A current multiplexing system comprising:

- a plurality of current multiplexers having at least one input and a plurality of outputs;
- a first and second alternating-current current source, each current source having a constant magnitude and frequency, wherein the second current source is complimentary to the first current source, and wherein each of the first and second current sources is coupled to an input of a current multiplexer;
- a plurality of current delivery circuits, each current delivery circuit having an input, a current output and a

30

voltage output, wherein each current delivery circuit input is coupled to an output of a current multiplexer;

- a plurality of electrodes, each of the plurality of electrodes being coupled to a current output of one of the plurality of current delivery circuits; and
- at least one voltage multiplexer having a plurality of inputs and at least one output:

wherein each current delivery circuit comprises:

- a current conveyor, having an input and an output, the current conveyor input being coupled to the current¹⁰ delivery circuit input and the current conveyor output being coupled to the current delivery circuit current output; and
- a first buffer having:
 - ¹⁵ a voltage buffer, having an input and an output, the voltage buffer input being coupled to the current delivery circuit current output and the voltage buffer output being coupled to the current delivery circuit voltage output;
 - a DC offset control circuit including an integrator coupled between the voltage buffer output and the current conveyor input; and,
 - a negative impedance circuit coupled between the voltage buffer output and the current conveyor ²⁵ output, the negative impedance circuit comprising:
 - a compensation impedance having a compensation impedance value, a first terminal and a second terminal, the first terminal coupled to $_{30}$ the current conveyor output; and
 - a compensation amplifier having a compensation amplifier input, a compensation amplifier output and a gain, the compensation amplifier input coupled to the voltage buffer output, the compensation amplifier output coupled to the second terminal of the compensation impedance to provide a compensation current to flow through the impedance, and the gain and the compensation impedance values are selected based on the parasitic impedance value so that the compensation current has a magnitude substantially equal to the leakage current magnitude.

18. Apparatus for electrically connecting measurement apparatus to a biological subject, the apparatus including a 45 signal delivery circuit including:

- a first buffer having:
 - a first buffer input for receiving a signal from a signal source; and,
 - a first buffer output for supplying a drive signal to an $_{50}$ electrode attached to the biological subject;
- a second buffer having:
 - a second buffer input coupled to the first buffer output; and,
 - a second buffer output for providing a sensed signal indicative of a signal at the electrode, to a sensor; and,
- a DC offset control circuit coupled between the second buffer output and the first buffer input.

19. Apparatus according to claim **18**, wherein the first $_{60}$ buffer is a current buffer, the drive signal being a drive current.

20. Apparatus according to claim **19**, wherein the current buffer is a current conveyor.

21. Apparatus according to claim **18**, wherein the second buffer is a voltage buffer, the sensed signal being a voltage signal indicative of a voltage at the electrode.

22. Apparatus according to claim 18, wherein the second buffer is a current buffer, the sensed signal being indicative of a drive current.

23. Apparatus for use in performing impedance measurements on a subject, wherein the apparatus includes:

- a signal source for applying a signal to a subject;
- a sensor for sensing signals from the subject;
- a processing system for controlling the signal source and receiving signals from the sensor; and,
- at least one signal delivery circuit, including:
 - a first buffer having:
 - a first buffer input for receiving a signal from a signal source; and,
 - a first buffer output for supplying a drive signal to an electrode attached to the biological subject;
 - a second buffer having:
 - a second buffer input coupled to the first buffer output; and,
 - a second buffer output for providing a sensed signal indicative of a signal at the electrode, to a sensor; and
 - a DC offset control circuit coupled between the second buffer output and the first buffer input.

24. Apparatus according to claim 23, wherein the apparatus includes:

- a first signal delivery circuit coupled to a drive electrode attached to the biological subject, the first signal delivery circuit including:
 - a first buffer having:
 - a first buffer input for receiving a signal from the signal source; and,
 - a first buffer output for supplying the drive signal to the drive electrode;
 - a second buffer having:
 - a second buffer input coupled to the first buffer output; and,
 - a second buffer output for providing a sensed signal indicative of the drive signal at the drive electrode; and
 - a DC offset control circuit coupled between the second buffer output and the first buffer input; and
 - a second signal delivery circuit coupled to a sense electrode attached to the biological subject, the second signal delivery circuit including:
 - a first buffer having:
 - a first buffer input coupled to ground; and,
 - a first buffer output coupled to the sense electrode; and,
 - a second buffer having:
 - a second buffer input coupled to the first buffer output;
 - a second buffer output for providing a sensed signal indicative of a sensed voltage at the sense electrode; and
 - a DC offset control circuit coupled between the second buffer output and the first buffer input.

* * * * *