



US009072495B2

(12) **United States Patent**  
**Specht**(10) **Patent No.:** **US 9,072,495 B2**  
(45) **Date of Patent:** **\*Jul. 7, 2015**(54) **METHOD AND APPARATUS TO PRODUCE  
ULTRASONIC IMAGES USING MULTIPLE  
APERTURES**(71) Applicant: **Maui Imaging, Inc.**, Sunnyvale, CA  
(US)  
(72) Inventor: **Donald F. Specht**, Los Altos, CA (US)  
(73) Assignee: **MAUI IMAGING, INC.**, Sunnyvale,  
CA (US)(\*) Notice: Subject to any disclaimer, the term of this  
patent is extended or adjusted under 35  
U.S.C. 154(b) by 0 days.This patent is subject to a terminal dis-  
claimer.(21) Appl. No.: **14/157,257**(22) Filed: **Jan. 16, 2014**(65) **Prior Publication Data**

US 2014/0135626 A1 May 15, 2014

**Related U.S. Application Data**(63) Continuation of application No. 13/632,929, filed on  
Oct. 1, 2012, now Pat. No. 8,684,936, which is a  
continuation of application No. 13/215,966, filed on  
Aug. 23, 2011, now Pat. No. 8,277,383, which is a  
(Continued)(51) **Int. Cl.**  
**A61B 8/14** (2006.01)  
**A61B 8/08** (2006.01)  
(Continued)(52) **U.S. Cl.**  
CPC ..... **A61B 8/5253** (2013.01); **A61B 8/42**  
(2013.01); **A61B 8/4209** (2013.01);  
(Continued)(58) **Field of Classification Search**CPC ..... A61B 8/145; A61B 8/14; A61B 8/42;  
A61B 8/4455; A61B 8/4209; A61B 8/4494;A61B 8/5253; A61B 8/4281; A61B 8/483;  
A61B 8/543; A61B 5/725; A61B 8/085;  
A61B 8/5207; G01S 7/52046; G01S 7/5205;  
G01S 15/8977USPC ..... 600/437-469; 382/128-132  
See application file for complete search history.(56) **References Cited**

## U.S. PATENT DOCUMENTS

3,174,286 A 3/1965 Erickson  
3,895,381 A 7/1975 Kock

(Continued)

## FOREIGN PATENT DOCUMENTS

EP 1949856 A1 7/2008  
EP 2187813 A1 5/2010

(Continued)

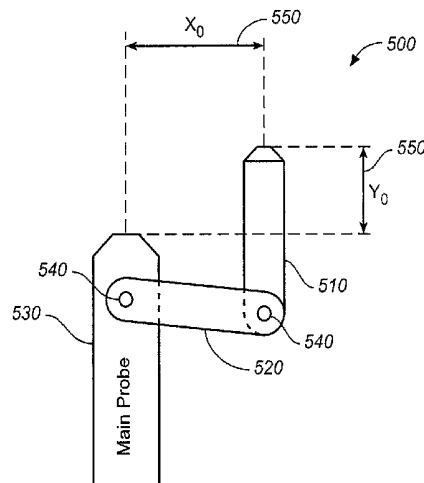
## OTHER PUBLICATIONS

Specht et al.; U.S. Appl. No. 14/279,052 entitled "Ultrasound imag-  
ing using apparent point-source transmit transducer," filed May 15,  
2014.

(Continued)

*Primary Examiner* — Sanjay Cattungal(74) *Attorney, Agent, or Firm* — Shay Glenn LLP(57) **ABSTRACT**

A combination of an ultrasonic scanner and an omnidirectional receive transducer for producing a two-dimensional image from received echoes is described. Two-dimensional images with different noise components can be constructed from the echoes received by additional transducers. These can be combined to produce images with better signal to noise ratios and lateral resolution. Also disclosed is a method based on information content to compensate for the different delays for different paths through intervening tissue is described. The disclosed techniques have broad application in medical imaging but are ideally suited to multi-aperture cardiac imaging using two or more intercostal spaces. Since lateral resolution is determined primarily by the aperture defined by the end elements, it is not necessary to fill the entire aperture with equally spaced elements. Multiple slices using these methods can be combined to form three-dimensional images.

**8 Claims, 11 Drawing Sheets**

**Related U.S. Application Data**

- continuation of application No. 11/865,501, filed on Oct. 1, 2007, now Pat. No. 8,007,439.
- (60) Provisional application No. 60/862,951, filed on Oct. 25, 2006, provisional application No. 60/940,261, filed on May 25, 2007.
- (51) **Int. Cl.**  
*A61B 8/00* (2006.01)  
*G01S 7/52* (2006.01)  
*G01S 15/89* (2006.01)  
*A61B 5/00* (2006.01)
- (52) **U.S. Cl.**  
 CPC ..... *A61B 8/4281* (2013.01); *A61B 8/4455* (2013.01); *A61B 8/483* (2013.01); *A61B 8/543* (2013.01); *G01S 7/52046* (2013.01); *G01S 7/5205* (2013.01); *G01S 15/8977* (2013.01); *A61B 5/725* (2013.01); *A61B 8/085* (2013.01); *A61B 8/145* (2013.01); *A61B 8/4494* (2013.01); *A61B 8/5207* (2013.01)

**(56) References Cited**

## U.S. PATENT DOCUMENTS

4,055,988 A	11/1977	Dutton	5,570,691 A	11/1996	Wright et al.
4,072,922 A	2/1978	Taner et al.	5,581,517 A	12/1996	Gee et al.
4,097,835 A	6/1978	Green	5,625,149 A	4/1997	Gururaja et al.
4,105,018 A	8/1978	Greenleaf et al.	5,628,320 A	5/1997	Teo
4,259,733 A	3/1981	Taner et al.	5,673,697 A	10/1997	Bryan et al.
4,271,842 A	6/1981	Specht et al.	5,675,550 A	10/1997	Ekhaus
4,325,257 A	4/1982	Kino et al.	5,720,291 A	2/1998	Schwartz
4,333,474 A	6/1982	Nigam	5,720,708 A	2/1998	Lu et al.
4,339,952 A	7/1982	Foster	5,744,898 A	4/1998	Smith et al.
4,452,084 A	6/1984	Taenzer	5,769,079 A	6/1998	Hossack
4,501,279 A	2/1985	Seo	5,784,334 A	7/1998	Sena et al.
4,539,847 A	9/1985	Paap	5,785,654 A	7/1998	Inuma et al.
4,566,459 A	1/1986	Umemura et al.	5,795,297 A	8/1998	Daigle
4,567,768 A	2/1986	Satoh et al.	5,797,845 A	8/1998	Barabash et al.
4,662,222 A	5/1987	Johnson	5,798,459 A	8/1998	Ohba et al.
4,669,482 A	6/1987	Ophir	5,838,564 A	11/1998	Bahorich et al.
4,682,497 A	7/1987	Sasaki	5,850,622 A	12/1998	Vassiliou et al.
4,781,199 A	11/1988	Hirama et al.	5,862,100 A	1/1999	VerWest
4,817,434 A	4/1989	Anderson	5,870,691 A	2/1999	Partyka et al.
4,831,601 A	5/1989	Breimesser et al.	5,876,342 A	3/1999	Chen et al.
4,893,284 A	1/1990	Magrane	5,891,038 A	4/1999	Seyed-Bolorforosh et al.
4,893,628 A	1/1990	Angelsen	5,892,732 A	4/1999	Gersztenkorn
5,050,588 A	9/1991	Grey et al.	5,916,169 A	6/1999	Hanafy et al.
5,141,738 A	8/1992	Rasor et al.	5,919,139 A	7/1999	Lin
5,161,536 A	11/1992	Vilkomerson et al.	5,920,285 A	7/1999	Benjamin
5,197,475 A	3/1993	Antich et al.	5,930,730 A	7/1999	Marfurt et al.
5,226,019 A	7/1993	Bahorich	5,940,778 A	8/1999	Marfurt et al.
5,230,339 A	7/1993	Charlebois	5,964,707 A	10/1999	Fenster et al.
5,269,309 A	12/1993	Fort et al.	5,969,661 A	10/1999	Benjamin
5,278,757 A	1/1994	Hector et al.	5,999,836 A	12/1999	Nelson et al.
5,293,871 A	3/1994	Reinstein et al.	6,007,499 A	12/1999	Martin et al.
5,299,576 A	4/1994	Shiba	6,013,032 A	1/2000	Savord
5,301,674 A	4/1994	Erikson et al.	6,014,473 A	1/2000	Hossack et al.
5,305,756 A	4/1994	Entrekin et al.	6,048,315 A	4/2000	Chiao et al.
5,339,282 A	8/1994	Kuhn et al.	6,049,509 A	4/2000	Sonneland et al.
5,340,510 A	8/1994	Bowen	6,050,943 A	4/2000	Slayton et al.
5,345,426 A	9/1994	Lipschutz	6,056,693 A	5/2000	Haider
5,355,888 A	10/1994	Kendall	6,058,074 A	5/2000	Swan et al.
5,409,010 A	4/1995	Beach et al.	6,077,224 A	6/2000	Lang et al.
5,442,462 A	8/1995	Guisin	6,092,026 A	7/2000	Bahorich et al.
5,503,152 A	4/1996	Oakley et al.	6,122,538 A	9/2000	Sliwa, Jr. et al.
5,515,853 A	5/1996	Smith et al.	6,123,670 A	9/2000	Mo
5,522,393 A	6/1996	Phillips et al.	6,129,672 A	10/2000	Seward et al.
5,526,815 A	6/1996	Granz et al.	6,135,960 A	10/2000	Holmberg
5,544,659 A	8/1996	Banjanin	6,138,075 A	10/2000	Yost
5,558,092 A	9/1996	Unger et al.	6,148,095 A	11/2000	Prause et al.
5,564,423 A	10/1996	Mele et al.	6,162,175 A	12/2000	Marian, Jr. et al.
5,568,812 A	10/1996	Murashita et al.	6,166,384 A	12/2000	Dentinger et al.
			6,166,853 A	12/2000	Sapia et al.
			6,193,665 B1	2/2001	Hall et al.
			6,196,739 B1	3/2001	Silverbrook
			6,200,266 B1	3/2001	Shokrollahi et al.
			6,210,335 B1	4/2001	Miller
			6,213,958 B1	4/2001	Winder
			6,221,019 B1	4/2001	Kantorovich
			6,231,511 B1	5/2001	Bae
			6,238,342 B1	5/2001	Feleppa et al.
			6,246,901 B1	6/2001	Benaron
			6,251,073 B1	6/2001	Imran et al.
			6,264,609 B1	7/2001	Herrington et al.
			6,266,551 B1	7/2001	Osadchy et al.
			6,278,949 B1	8/2001	Alam
			6,289,230 B1	9/2001	Chaiken et al.
			6,304,684 B1	10/2001	Niczyporuk et al.
			6,309,356 B1	10/2001	Ustuner et al.
			6,324,453 B1	11/2001	Breed et al.
			6,345,539 B1	2/2002	Rawes et al.
			6,361,500 B1	3/2002	Masters
			6,363,033 B1	3/2002	Cole et al.
			6,374,185 B1	4/2002	Taner et al.
			6,394,955 B1	5/2002	Perlit
			6,423,002 B1	7/2002	Hossack
			6,436,046 B1	8/2002	Napolitano et al.
			6,449,821 B1	9/2002	Sudol et al.
			6,450,965 B2	9/2002	Williams et al.
			6,468,216 B1	10/2002	Powers et al.
			6,471,650 B2	10/2002	Powers et al.
			6,475,150 B2	11/2002	Haddad
			6,480,790 B1	11/2002	Calvert et al.

(56)

## References Cited

## U.S. PATENT DOCUMENTS

6,487,502	B1	11/2002	Taner	7,824,337	B2	11/2010	Abe et al.
6,499,536	B1	12/2002	Ellingsen	7,833,163	B2	11/2010	Cai
6,508,768	B1	1/2003	Hall et al.	7,837,624	B1	11/2010	Hossack et al.
6,517,484	B1	2/2003	Wilk et al.	7,846,097	B2	12/2010	Jones et al.
6,526,163	B1	2/2003	Halmann et al.	7,850,613	B2	12/2010	Stribling
6,543,272	B1	4/2003	Vitek	7,862,508	B2	1/2011	Davies et al.
6,547,732	B2	4/2003	Jago	7,876,945	B2	1/2011	Lötjönen
6,551,246	B1	4/2003	Ustuner et al.	7,887,486	B2	2/2011	Ustuner et al.
6,565,510	B1	5/2003	Haider	7,901,358	B2	3/2011	Mehi et al.
6,585,647	B1	7/2003	Winder	7,914,451	B2	3/2011	Davies
6,604,421	B1	8/2003	Li	7,919,906	B2	4/2011	Cerofolini
6,614,560	B1	9/2003	Silverbrook	7,926,350	B2	4/2011	Kröning et al.
6,620,101	B2	9/2003	Azzam et al.	7,927,280	B2	4/2011	Davidson
6,652,461	B1	11/2003	Levkovitz	7,972,271	B2	7/2011	Johnson et al.
6,668,654	B2	12/2003	Dubois et al.	7,984,637	B2	7/2011	Ao et al.
6,672,165	B2	1/2004	Rather et al.	7,984,651	B2	7/2011	Randall et al.
6,681,185	B1	1/2004	Young et al.	8,002,705	B1	8/2011	Napolitano et al.
6,690,816	B2	2/2004	Aylward et al.	8,007,439	B2	8/2011	Specht
6,692,450	B1	2/2004	Coleman	8,057,392	B2	11/2011	Hossack et al.
6,695,778	B2	2/2004	Golland et al.	8,057,393	B2	11/2011	Yao et al.
6,702,745	B1	3/2004	Smythe	8,079,263	B2	12/2011	Randall et al.
6,719,693	B2	4/2004	Richard	8,079,956	B2	12/2011	Azuma et al.
6,728,567	B2	4/2004	Rather et al.	8,088,067	B2	1/2012	Vortman et al.
6,752,762	B1	6/2004	DeJong et al.	8,088,068	B2	1/2012	Yao et al.
6,755,787	B2	6/2004	Hossack et al.	8,088,071	B2	1/2012	Hwang et al.
6,780,152	B2	8/2004	Ustuner et al.	8,105,239	B2	1/2012	Specht
6,790,182	B2	9/2004	Eck et al.	8,157,737	B2	4/2012	Zhang et al.
6,837,853	B2	1/2005	Marian	8,202,219	B2	6/2012	Luo et al.
6,843,770	B2	1/2005	Sumanaweera	8,277,383	B2	10/2012	Specht
6,847,737	B1	1/2005	Kouri et al.	8,412,307	B2	4/2013	Willis et al.
6,854,332	B2	2/2005	Alleyne	8,419,642	B2	4/2013	Sandrin et al.
6,932,767	B2	8/2005	Landry et al.	8,473,239	B2	6/2013	Specht et al.
7,033,320	B2	4/2006	Von Behren et al.	8,602,993	B2	12/2013	Specht et al.
7,087,023	B2	8/2006	Daft et al.	8,672,846	B2	3/2014	Napolitano et al.
7,104,956	B1	9/2006	Christopher	2002/0035864	A1	3/2002	Paltieli et al.
7,217,243	B2	5/2007	Takeuchi	2002/0087071	A1	7/2002	Schmitz et al.
7,221,867	B2	5/2007	Silverbrook	2002/0111568	A1	8/2002	Bukshpan
7,231,072	B2	6/2007	Yamano et al.	2003/0028111	A1	2/2003	Vaezy et al.
7,269,299	B2	9/2007	Schroeder	2003/0040669	A1	2/2003	Grass et al.
7,283,652	B2	10/2007	Mendonca et al.	2003/0228053	A1	12/2003	Li et al.
7,285,094	B2	10/2007	Nohara et al.	2004/0054283	A1	3/2004	Corey et al.
7,313,053	B2	12/2007	Wodnicki	2004/0068184	A1	4/2004	Trahey et al.
7,366,704	B2	4/2008	Reading et al.	2004/0100163	A1	5/2004	Baumgartner et al.
7,402,136	B2	7/2008	Hossack et al.	2004/0111028	A1	6/2004	Abe et al.
7,410,469	B1	8/2008	Talish et al.	2004/0122313	A1	6/2004	Moore et al.
7,415,880	B2	8/2008	Renzel	2004/0122322	A1	6/2004	Moore et al.
7,443,765	B2	10/2008	Thomenius et al.	2004/0127793	A1	7/2004	Mendlein et al.
7,444,875	B1	11/2008	Wu et al.	2004/0138565	A1	7/2004	Trucco
7,447,535	B2	11/2008	Lavi	2004/0144176	A1	7/2004	Yoden
7,448,998	B2	11/2008	Robinson	2004/0236217	A1	11/2004	Cerwin et al.
7,466,848	B2	12/2008	Metaxas et al.	2004/0236223	A1	11/2004	Barnes et al.
7,469,096	B2	12/2008	Silverbrook	2005/0004449	A1	1/2005	Mitschke et al.
7,474,778	B2	1/2009	Shinomura et al.	2005/0053305	A1	3/2005	Li et al.
7,481,577	B2	1/2009	Ramamurthy et al.	2005/0054910	A1	3/2005	Tremblay et al.
7,491,171	B2	2/2009	Barthe et al.	2005/0090743	A1	4/2005	Kawashima et al.
7,497,828	B1	3/2009	Wilk et al.	2005/0090745	A1	4/2005	Steen
7,497,830	B2	3/2009	Li	2005/0111846	A1	5/2005	Steinbacher et al.
7,510,529	B2	3/2009	Chou et al.	2005/0113694	A1	5/2005	Haugen et al.
7,514,851	B2	4/2009	Wilser et al.	2005/0124883	A1	6/2005	Hunt
7,549,962	B2	6/2009	Dreschel et al.	2005/0131300	A1	6/2005	Bakircioglu et al.
7,574,026	B2	8/2009	Rasche et al.	2005/0147297	A1	7/2005	McLaughlin et al.
7,625,343	B2	12/2009	Cao et al.	2005/0165312	A1	7/2005	Knowles et al.
7,637,869	B2	12/2009	Sudol	2005/0203404	A1	9/2005	Freiburger
7,668,583	B2	2/2010	Fegert et al.	2005/0215883	A1	9/2005	Hundley et al.
7,674,228	B2	3/2010	Williams et al.	2005/0240125	A1	10/2005	Makin et al.
7,682,311	B2	3/2010	Simopoulos et al.	2005/0281447	A1	12/2005	Moreau-Gobard et al.
7,699,776	B2	4/2010	Walker et al.	2005/0288588	A1	12/2005	Weber et al.
7,722,541	B2	5/2010	Cai	2006/0062447	A1	3/2006	Rinck et al.
7,744,532	B2	6/2010	Ustuner et al.	2006/0074313	A1	4/2006	Slayton et al.
7,750,311	B2	7/2010	Daghighian	2006/0074315	A1	4/2006	Liang et al.
7,785,260	B2	8/2010	Umemura et al.	2006/0074320	A1	4/2006	Yoo et al.
7,787,680	B2	8/2010	Ahn et al.	2006/0079759	A1	4/2006	Vaillant et al.
7,806,828	B2	10/2010	Stringer	2006/0079778	A1	4/2006	Mo et al.
7,819,810	B2	10/2010	Stringer et al.	2006/0079782	A1	4/2006	Beach et al.
7,822,250	B2	10/2010	Yao et al.	2006/0094962	A1	5/2006	Clark
				2006/0122506	A1	6/2006	Davies et al.
				2006/0173327	A1	8/2006	Kim
				2006/0262291	A1	11/2006	Hess et al.
				2007/0016022	A1	1/2007	Blalock et al.

(56)

## References Cited

## U.S. PATENT DOCUMENTS

2007/0016044 A1 1/2007 Blalock et al.  
 2007/0036414 A1 2/2007 Georgescu et al.  
 2007/0055155 A1 3/2007 Owen et al.  
 2007/0078345 A1 4/2007 Mo et al.  
 2007/0088213 A1 4/2007 Poland  
 2007/0138157 A1 6/2007 Dane et al.  
 2007/0161898 A1 7/2007 Hao et al.  
 2007/0161904 A1 7/2007 Urbano  
 2007/0167752 A1 7/2007 Proulx et al.  
 2007/0167824 A1 7/2007 Lee et al.  
 2007/0232914 A1 10/2007 Chen et al.  
 2007/0238985 A1 10/2007 Smith et al.  
 2007/0242567 A1 10/2007 Daft et al.  
 2008/0110261 A1 5/2008 Randall et al.  
 2008/0110263 A1 5/2008 Klessel et al.  
 2008/0112265 A1 5/2008 Urbano et al.  
 2008/0114241 A1 5/2008 Randall et al.  
 2008/0114245 A1 5/2008 Randall et al.  
 2008/0114246 A1 5/2008 Randall et al.  
 2008/0114247 A1 5/2008 Urbano et al.  
 2008/0114248 A1 5/2008 Urbano et al.  
 2008/0114249 A1 5/2008 Randall et al.  
 2008/0114250 A1 5/2008 Urbano et al.  
 2008/0114251 A1 5/2008 Weymer et al.  
 2008/0114252 A1 5/2008 Randall et al.  
 2008/0114253 A1 5/2008 Randall et al.  
 2008/0114255 A1 5/2008 Schwartz et al.  
 2008/0125659 A1 5/2008 Wilser et al.  
 2008/0181479 A1 7/2008 Yang et al.  
 2008/0183075 A1 7/2008 Govari et al.  
 2008/0188747 A1 8/2008 Randall et al.  
 2008/0188750 A1 8/2008 Randall et al.  
 2008/0194957 A1 8/2008 Hctor et al.  
 2008/0194958 A1 8/2008 Lee et al.  
 2008/0194959 A1 8/2008 Wang et al.  
 2008/0208061 A1 8/2008 Halmann  
 2008/0242996 A1 10/2008 Hall et al.  
 2008/0249408 A1 10/2008 Palmeri et al.  
 2008/0255452 A1 10/2008 Entrekin  
 2008/0269604 A1 10/2008 Bector et al.  
 2008/0269613 A1 10/2008 Summers et al.  
 2008/0275344 A1 11/2008 Glide-Hurst et al.  
 2008/0285819 A1 11/2008 Konofagou et al.  
 2008/0287787 A1 11/2008 Sauer et al.  
 2008/0294045 A1 11/2008 Ellington et al.  
 2008/0294050 A1 11/2008 Shinomura et al.  
 2008/0294052 A1 11/2008 Wilser et al.  
 2008/0306382 A1 12/2008 Guracar et al.  
 2008/0306386 A1 12/2008 Baba et al.  
 2008/0319317 A1 12/2008 Kamiyama et al.  
 2009/0010459 A1 1/2009 Garbini et al.  
 2009/0012393 A1 1/2009 Choi  
 2009/0016163 A1 1/2009 Freeman et al.  
 2009/0018445 A1 1/2009 Schers et al.  
 2009/0024039 A1 1/2009 Wang et al.  
 2009/0036780 A1 2/2009 Abraham  
 2009/0043206 A1 2/2009 Towfiq et al.  
 2009/0048519 A1 2/2009 Hossack et al.  
 2009/0069681 A1 3/2009 Lundberg et al.  
 2009/0069686 A1 3/2009 Daft et al.  
 2009/0069692 A1 3/2009 Cooley et al.  
 2009/0112095 A1 4/2009 Daigle  
 2009/0182237 A1 7/2009 Angelsen et al.  
 2009/0208080 A1 8/2009 Grau et al.  
 2009/0306510 A1 12/2009 Hashiba et al.  
 2010/0016725 A1 1/2010 Thiele  
 2010/0121193 A1 5/2010 Fukukita et al.  
 2010/0121196 A1 5/2010 Hwang et al.  
 2010/0168578 A1 7/2010 Garson, Jr. et al.  
 2010/0217124 A1 8/2010 Cooley  
 2010/0256488 A1 10/2010 Kim et al.  
 2010/0262013 A1 10/2010 Smith et al.  
 2011/0098565 A1 4/2011 Masuzawa  
 2011/0112404 A1 5/2011 Gourevitch  
 2011/0125017 A1 5/2011 Ramamurthy et al.

2011/0201933 A1 8/2011 Specht et al.  
 2012/0036934 A1 2/2012 Kröning et al.  
 2012/0057428 A1 3/2012 Specht et al.  
 2012/0095343 A1 4/2012 Smith et al.  
 2012/0095347 A1 4/2012 Adam et al.  
 2012/0116226 A1 5/2012 Specht  
 2013/0035595 A1 2/2013 Specht  
 2013/0144166 A1 6/2013 Specht et al.  
 2013/0172743 A1 7/2013 Brewer et al.  
 2013/0218012 A1 8/2013 Specht et al.  
 2013/0247350 A1 9/2013 Specht et al.  
 2013/0253325 A1 9/2013 Call et al.  
 2013/0261463 A1 10/2013 Chiang et al.  
 2014/0043933 A1 2/2014 Belevich et al.

## FOREIGN PATENT DOCUMENTS

EP 2198785 A1 6/2010  
 EP 1757955 B1 11/2010  
 EP 1979739 10/2011  
 EP 1840594 B1 6/2012  
 EP 1850743 B1 12/2012  
 EP 1594404 B1 9/2013  
 EP 2026280 B1 10/2013  
 FR 2851662 A1 8/2004  
 JP S49-11189 A 1/1974  
 JP S54-44375 4/1979  
 JP S55-103839 A 8/1980  
 JP S59-174151 A 10/1984  
 JP S60-13109 1/1985  
 JP S60-68836 A 4/1985  
 JP 4-67856 3/1992  
 JP 05-042138 A 2/1993  
 JP 7-051266 A 2/1995  
 JP 08-252253 10/1996  
 JP 9-103429 A 4/1997  
 JP 9-201361 A 8/1997  
 JP 10-216128 A 8/1998  
 JP 11-089833 A 4/1999  
 JP 2001-245884 A 9/2001  
 JP 2002-209894 A 7/2002  
 JP 2002-253549 9/2002  
 JP 2004-215987 8/2004  
 JP 2004-337457 12/2004  
 JP 2005-523792 8/2005  
 JP 2005-526539 9/2005  
 JP 2008-122209 5/2008  
 JP 2008-513763 A 5/2008  
 JP 2008-259541 A 10/2008  
 WO WO 92/18054 A1 10/1992  
 WO WO 98/00719 A2 1/1998  
 WO WO02/084594 A2 10/2002  
 WO WO2005/009245 A1 2/2005  
 WO WO 2006/114735 A1 11/2006  
 WO WO 2007/127147 A2 11/2007

## OTHER PUBLICATIONS

Cristianini et al.; An Introduction to Support Vector Machines; Cambridge University Press; pp. 93-111; Mar. 2000.  
 Feigenbaum, Harvey, M.D.; Echocardiography; Lippincott Williams & Wilkins; Philadelphia; 5th Ed.; pp. 428, 484; Feb. 1994.  
 Haykin, Simon; Neural Networks: A Comprehensive Foundation (2nd Ed.); Prentice Hall; pp. 156-187; Jul. 16, 1998.  
 Kramb et al.; Considerations for using phased array ultrasonics in a fully automated inspection system. Review of Quantitative Nondestructive Evaluation, vol. 23, ed. D. O. Thompson and D. E. Chimenti, pp. 817-825, (year of publication is sufficiently earlier than the effective U.S. filing date and any foreign priority date) 2004.  
 Ledesma-Carbayo et al.; Spatio-temporal nonrigid registration for ultrasound cardiac motion estimation; IEEE Trans. on Medical Imaging; vol. 24; No. 9; Sep. 2005.  
 Leotta et al.; Quantitative three-dimensional echocardiography by rapid imaging . . . ; J American Society of Echocardiography; vol. 10; No. 8; pp. 830-839; Oct. 1997.  
 Morrison et al.; A probabilistic neural network based image segmentation network for magnetic resonance images; Proc. Conf. Neural Networks; Baltimore, MD; vol. 3; pp. 60-65; Jun. 1992.

(56)

**References Cited**

## OTHER PUBLICATIONS

Nadkarni et al.; Cardiac motion synchronization for 3D cardiac ultrasound imaging; Ph.D. Dissertation, University of Western Ontario; Jun. 2002.

Press et al.; Cubic spline interpolation; §3.3 in "Numerical Recipes in FORTRAN: The Art of Scientific Computing", 2nd Ed.; Cambridge, England; Cambridge University Press; pp. 107-110; Sep. 1992.

Sakas et al.; Preprocessing and volume rendering of 3D ultrasonic data; IEEE Computer Graphics and Applications; pp. 47-54, Jul. 1995.

Sapia et al.; Deconvolution of ultrasonic waveforms using an adaptive wiener filter; Review of Progress in Quantitative Nondestructive Evaluation; vol. 13A; Plenum Press; pp. 855-862; (year of publication is sufficiently earlier than the effective U.S. filed and any foreign priority date) 1994.

Sapia et al.; Ultrasound image deconvolution using adaptive inverse filtering; 12 IEEE Symposium on Computer-Based Medical Systems, CBMS, pp. 248-253; Jun. 1999.

Sapia, Mark Angelo; Multi-dimensional deconvolution of optical microscope and ultrasound imaging using adaptive least-mean-square (LMS) inverse filtering; Ph.D. Dissertation; University of Connecticut; Jan. 2000.

Smith et al.; High-speed ultrasound volumetric imaging system. 1. Transducer design and beam steering; IEEE Trans. Ultrason., Ferroelect., Freq. Contr.; vol. 38; pp. 100-108; Mar. 1991.

Specht et al.; Deconvolution techniques for digital longitudinal tomography; SPIE; vol. 454; presented at Application of Optical Instrumentation in Medicine XII; pp. 319-325; Jun. 1984.

Specht et al.; Experience with adaptive PNN and adaptive GRNN; Proc. IEEE International Joint Conf. on Neural Networks; vol. 2; pp. 1203-1208; Orlando, FL; Jun. 1994.

Specht, D.F.; A general regression neural network; IEEE Trans. on Neural Networks; vol. 2.; No. 6; Nov. 1991.

Specht, D.F.; Blind deconvolution of motion blur using LMS inverse filtering; Lockheed Independent Research (unpublished); Jun. 23, 1975.

Specht, D.F.; Enhancements to probabilistic neural networks; Proc. IEEE International Joint Conf. on Neural Networks; Baltimore, MD; Jun. 1992.

Specht, D.F.; GRNN with double clustering; Proc. IEEE International Joint Conf. Neural Networks; Vancouver, Canada; Jul. 16-21, 2006.

Specht, D.F.; Probabilistic neural networks; Pergamon Press; Neural Networks; vol. 3; pp. 109-118; Feb. 1990.

Von Ramm et al.; High-speed ultrasound volumetric imaging-System. 2. Parallel processing and image display; IEEE Trans. Ultrason., Ferroelect., Freq. Contr.; vol. 38; pp. 109-115; Mar. 1991.

Wells, P.N.T.; Biomedical ultrasonics; Academic Press; London, New York, San Francisco; pp. 124-125; Mar. 1977.

Widrow et al.; Adaptive signal processing; Prentice-Hall; Englewood Cliffs, NJ; pp. 99-116; Mar. 1985.

Call et al.; U.S. Appl. No. 13/971,689 entitled "Ultrasound Imaging System Memory Architecture," filed Aug. 20, 2013.

Specht et al.; U.S. Appl. No. 14/078,311 entitled "Imaging with Multiple Aperture Medical Ultrasound and Synchronization of Add-On Systems," filed Nov. 12, 2013.

Smith et al.; U.S. Appl. No. 14/210,015 entitled "Alignment of ultrasound transducer arrays and multiple aperture probe assembly," filed Mar. 13, 2014.

Smith et al.; U.S. Appl. No. 14/526,186 entitled "Universal multiple aperture medical ultrasound probe," filed Oct. 28, 2014.

Smith et al.; U.S. Appl. No. 14/595,083 entitled "Concave ultrasound transducers and 3D arrays," filed Jan. 12, 2015.

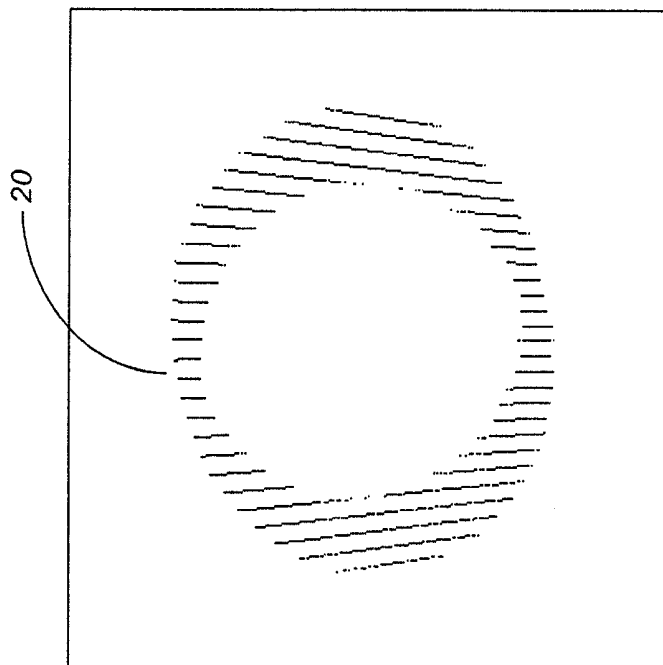
Hendee et al.; Medical Imaging Physics; Wiley-Liss, Inc. 4th Edition; Chap. 19-22; pp. 303-353; © 2002 (year of pub. sufficiently earlier than effective US filing date and any foreign priority date).

Wikipedia; Point cloud; 2 pages; Nov. 24, 2014; retrieved from the internet ([https://en.wikipedia.org/w/index.php?title=Point\\_cloud&oldid=472583138](https://en.wikipedia.org/w/index.php?title=Point_cloud&oldid=472583138)).

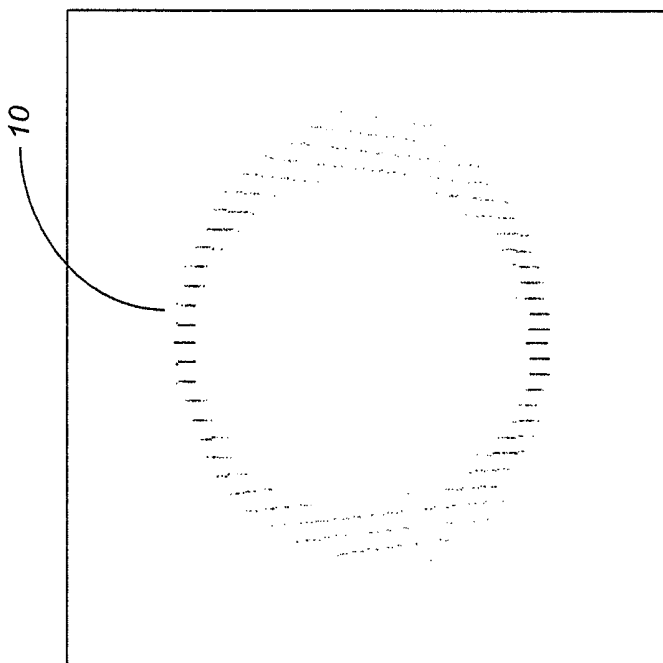
Li et al.; An efficient speckle tracking algorithm for ultrasonic imaging; 24; pp. 215-228; Oct. 1, 2002.

UCLA Academic Technology; SPSS learning module: How can I analyze a subset of my data; 6 pages; retrieved from the Internet ([http://www.ats.ucla.edu/stat/spss/modules/subset\\_analyze.htm](http://www.ats.ucla.edu/stat/spss/modules/subset_analyze.htm)) Nov. 26, 2001.

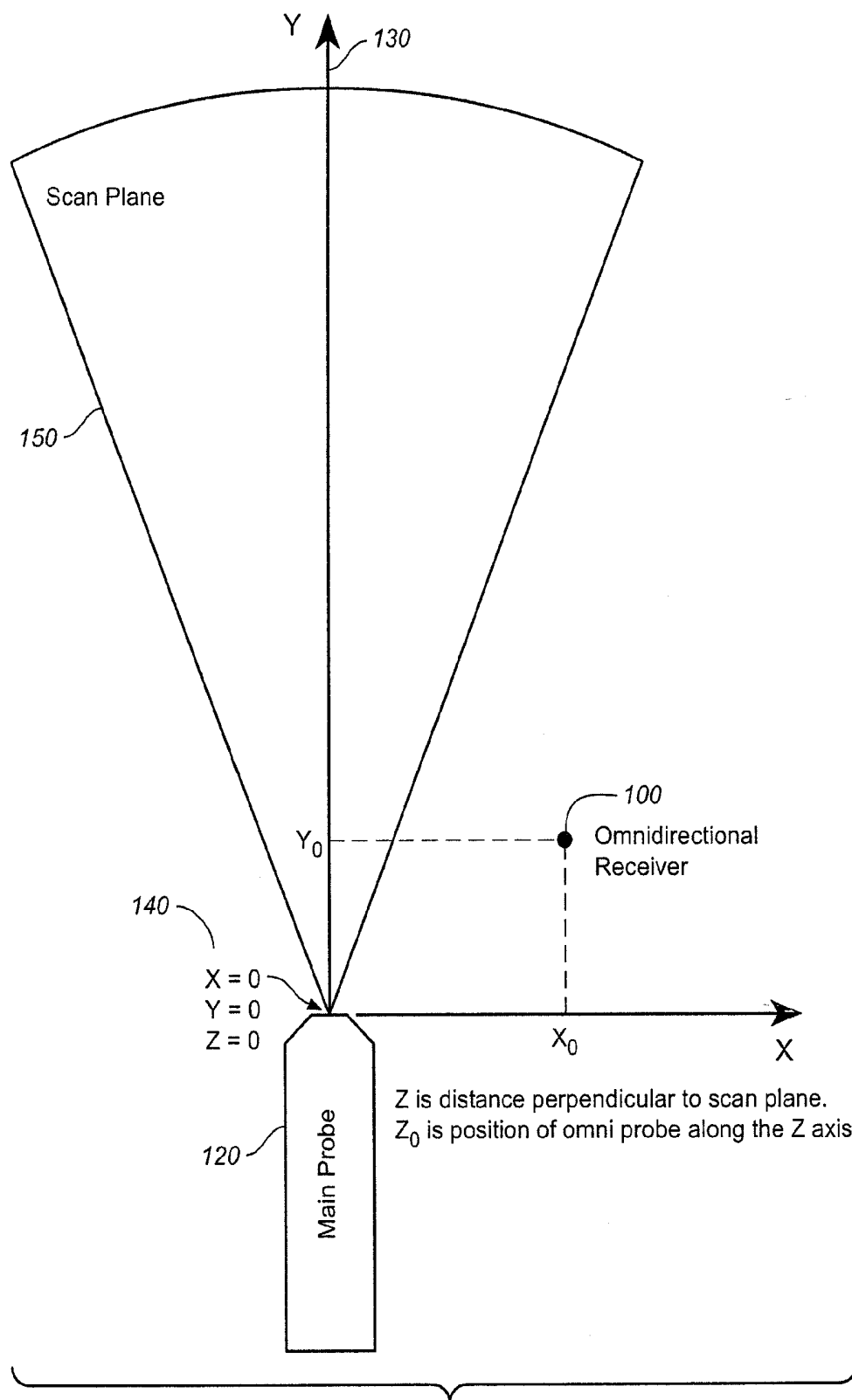
Wikipedia; Curve fitting; 5 pages; retrieved from the internet ([http://en.wikipedia.org/wiki/Curve\\_fitting](http://en.wikipedia.org/wiki/Curve_fitting)) Dec. 19, 2010.

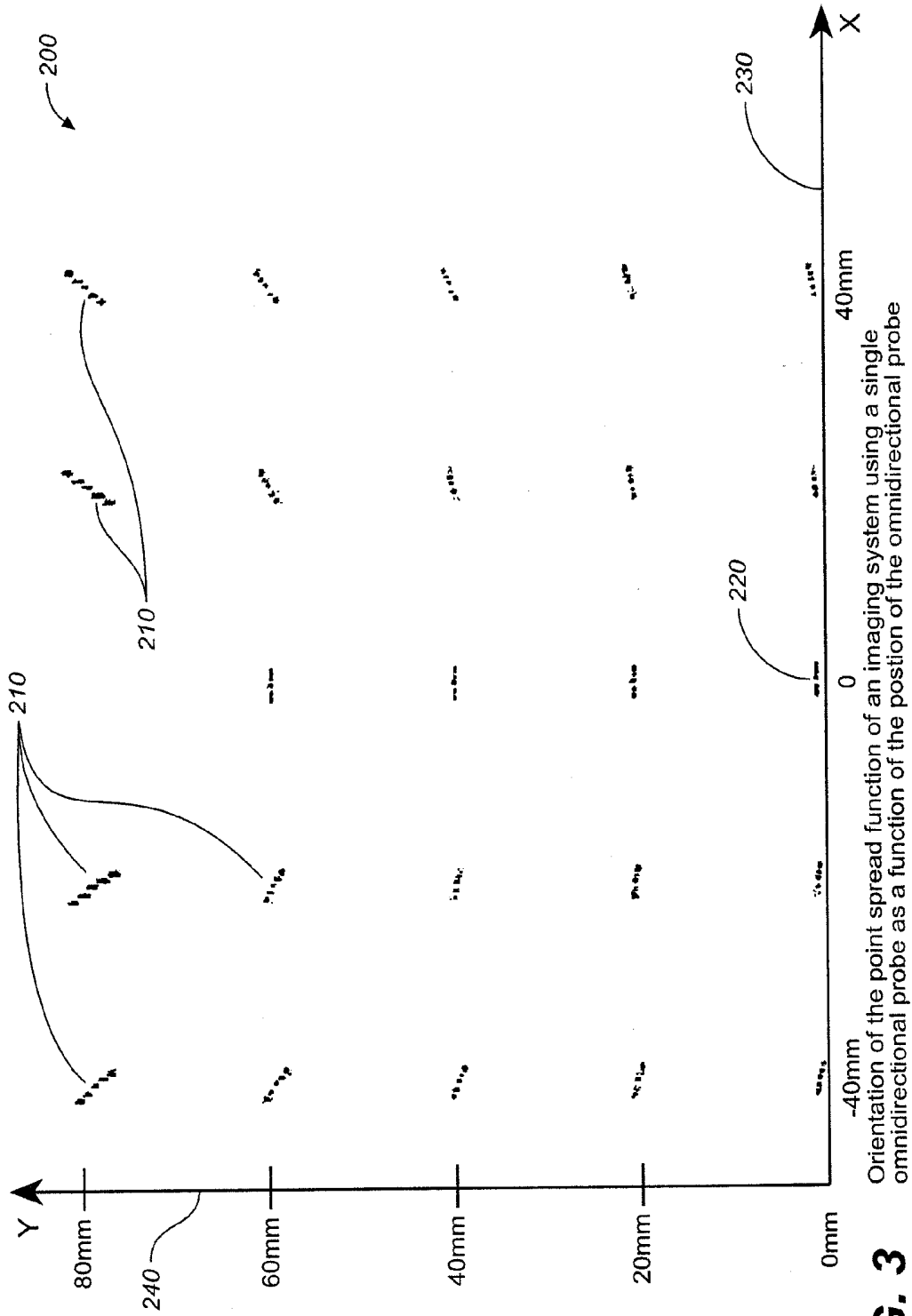


**FIG. 5**



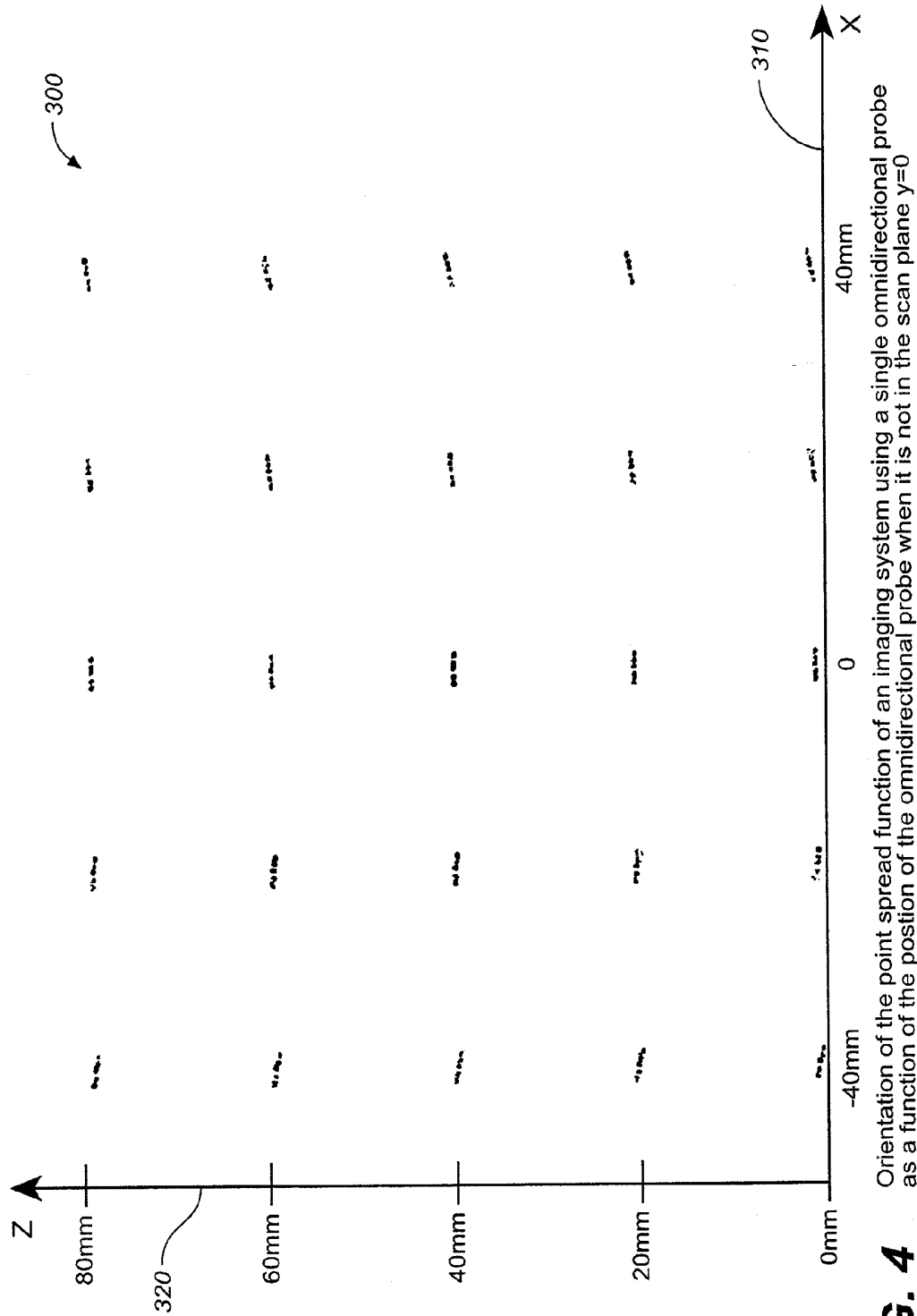
**FIG. 1**

**FIG. 2**

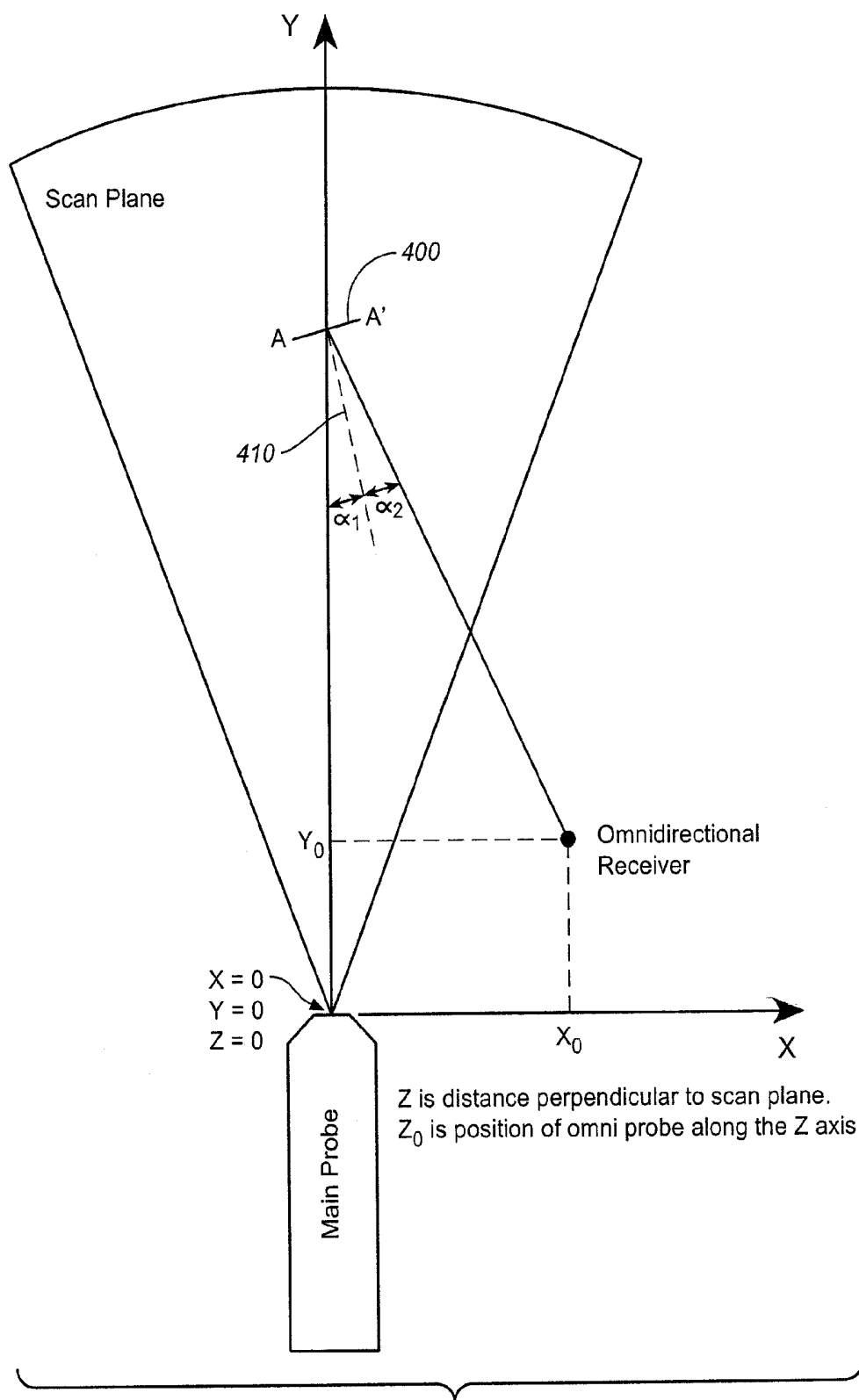


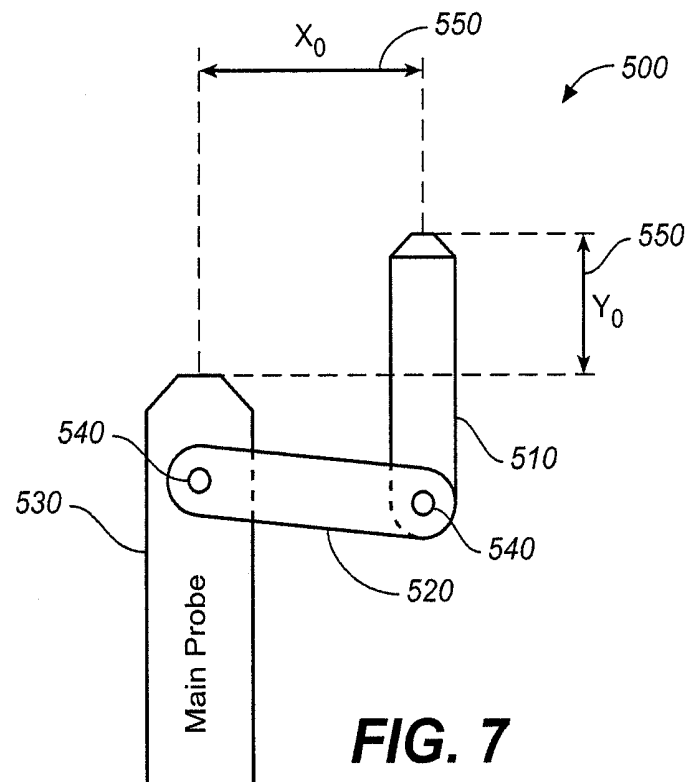
**FIG. 3**

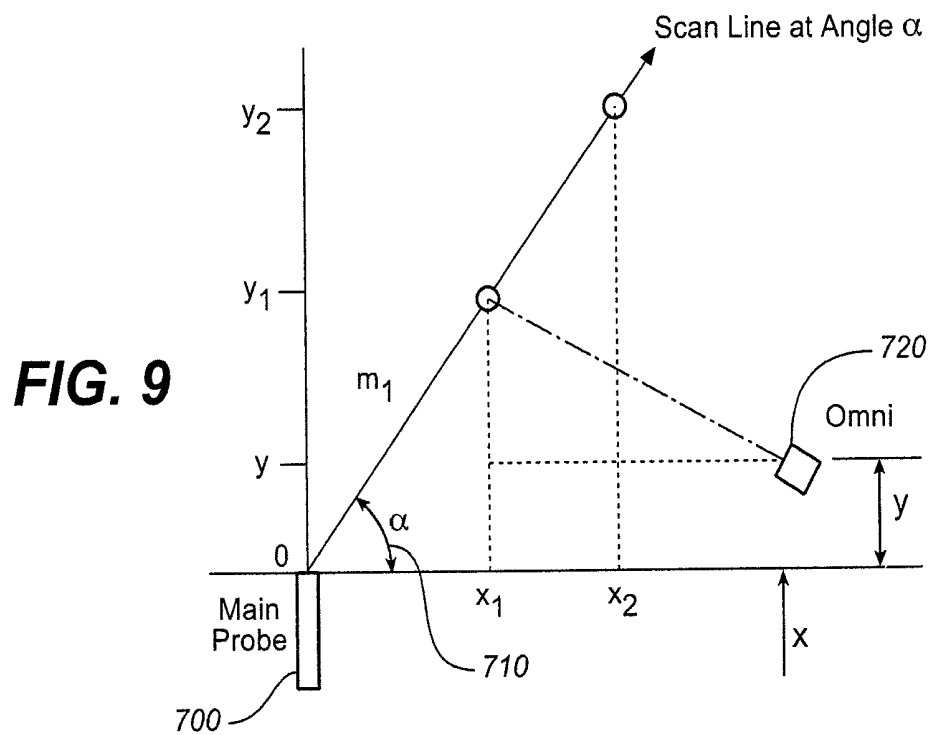
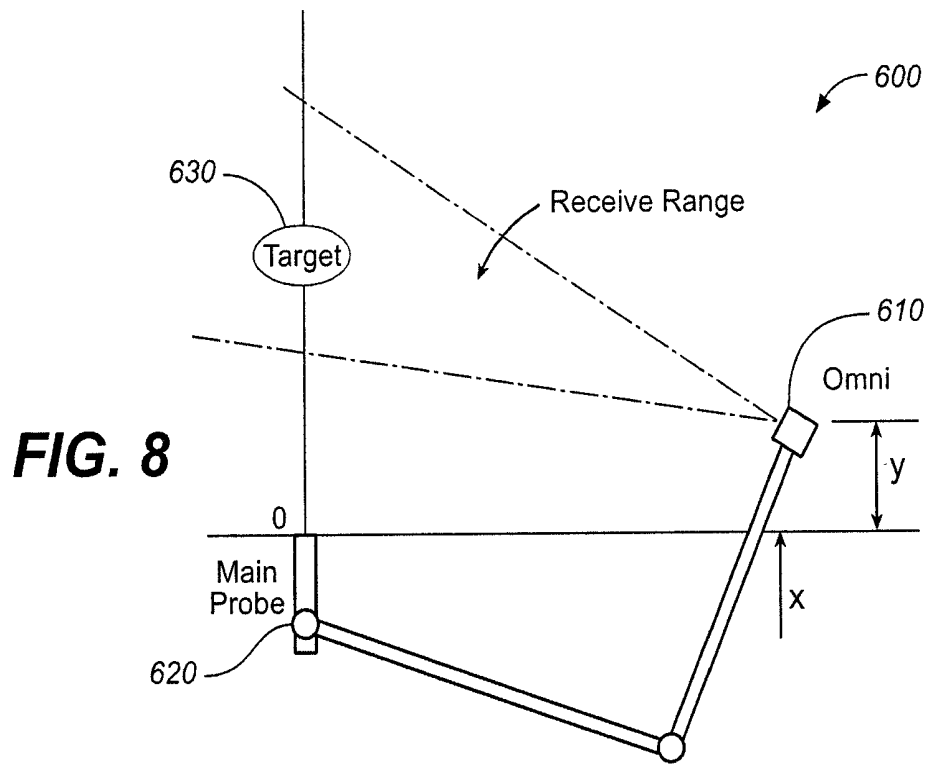


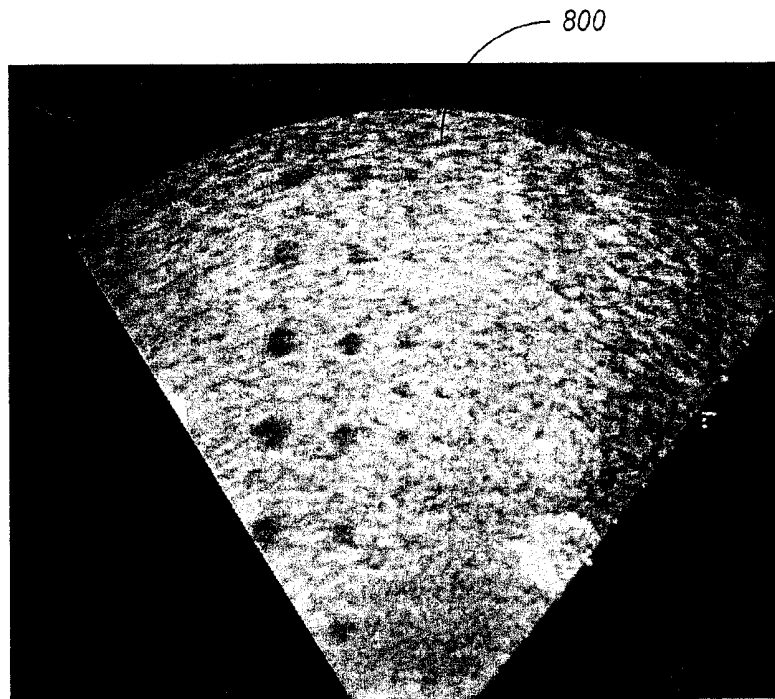


**FIG. 4** Orientation of the point spread function of an imaging system using a single omnidirectional probe as a function of the position of the omnidirectional probe when it is not in the scan plane  $y=0$

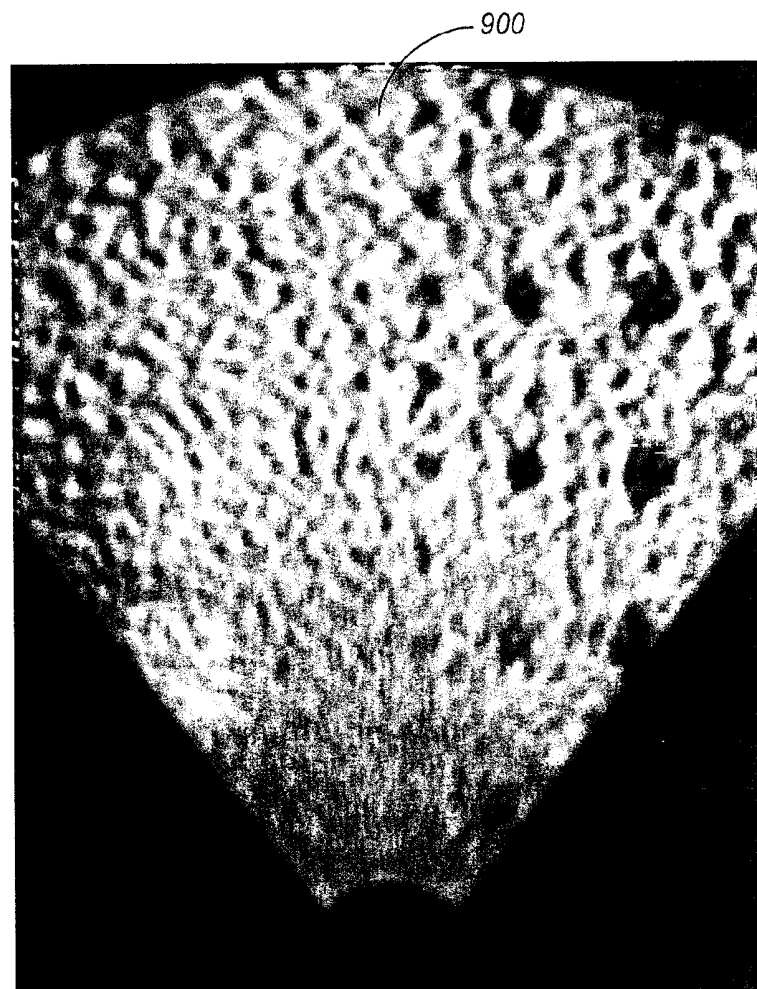




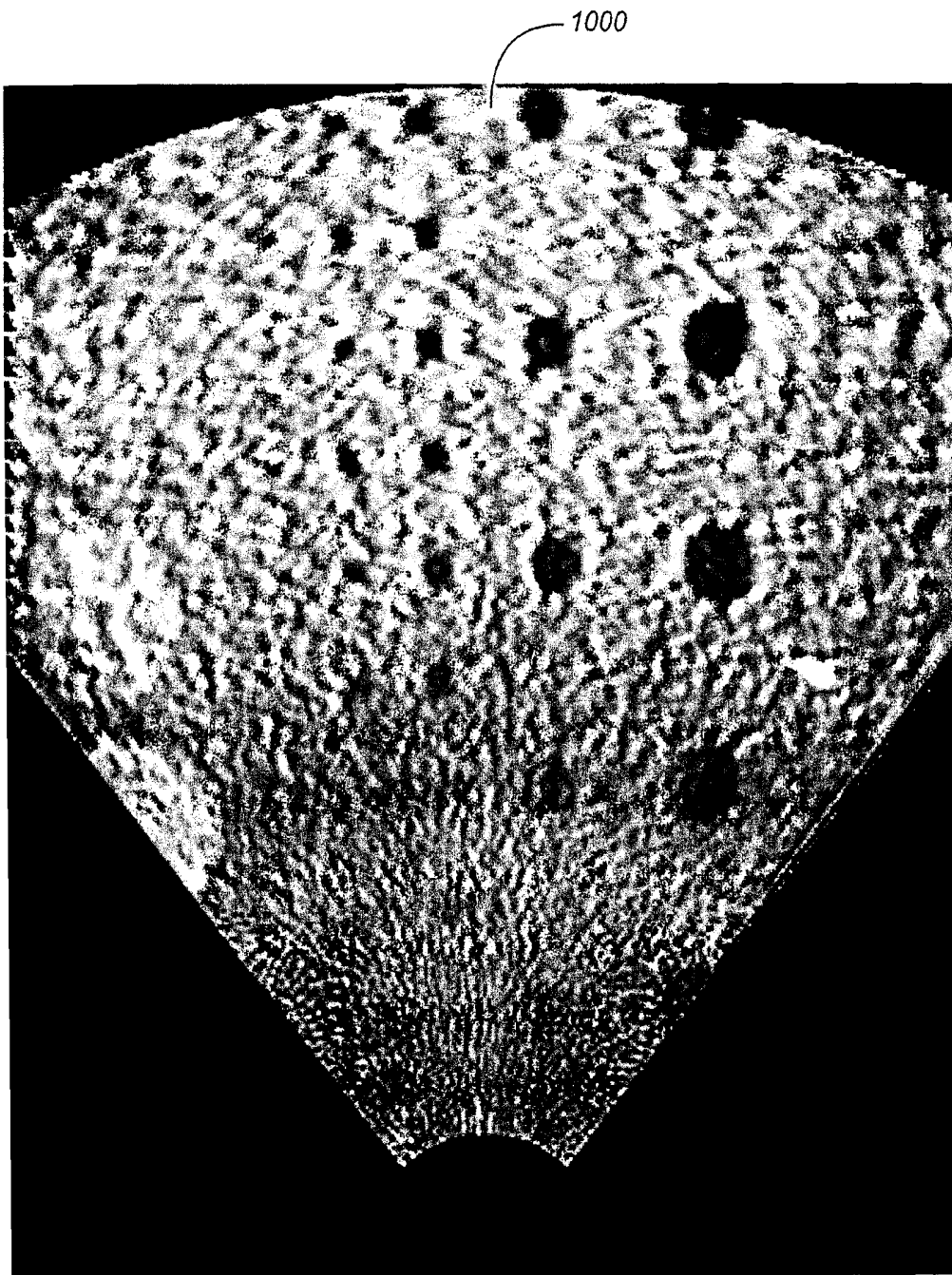




**FIG. 10A**



**FIG. 10B**



**FIG. 11**

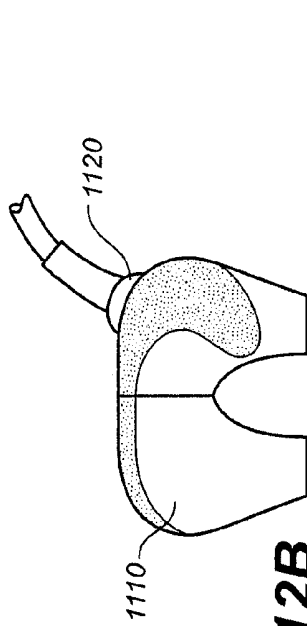


FIG. 12B

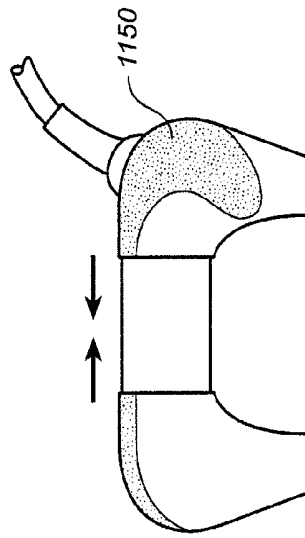


FIG. 12C

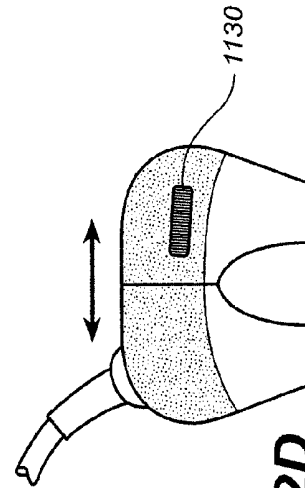


FIG. 12D

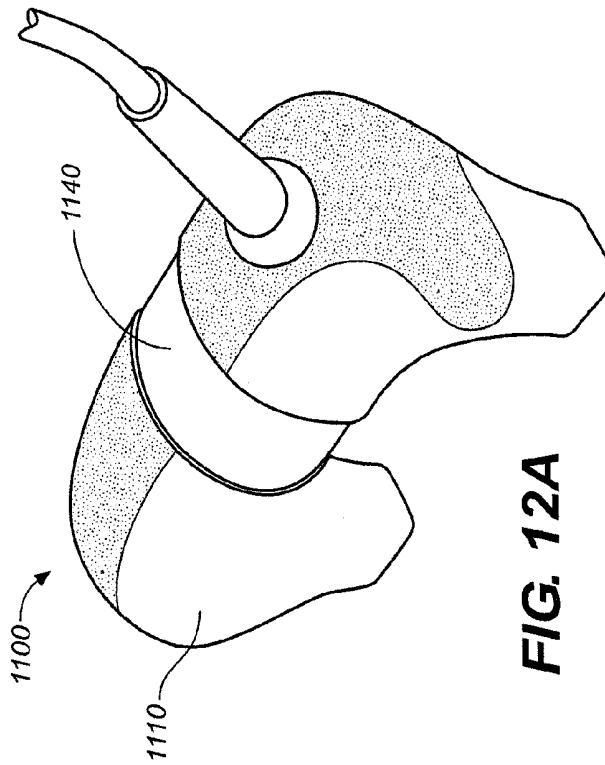


FIG. 12A

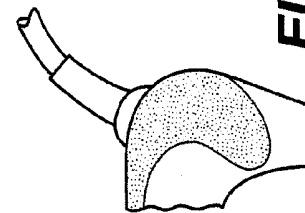
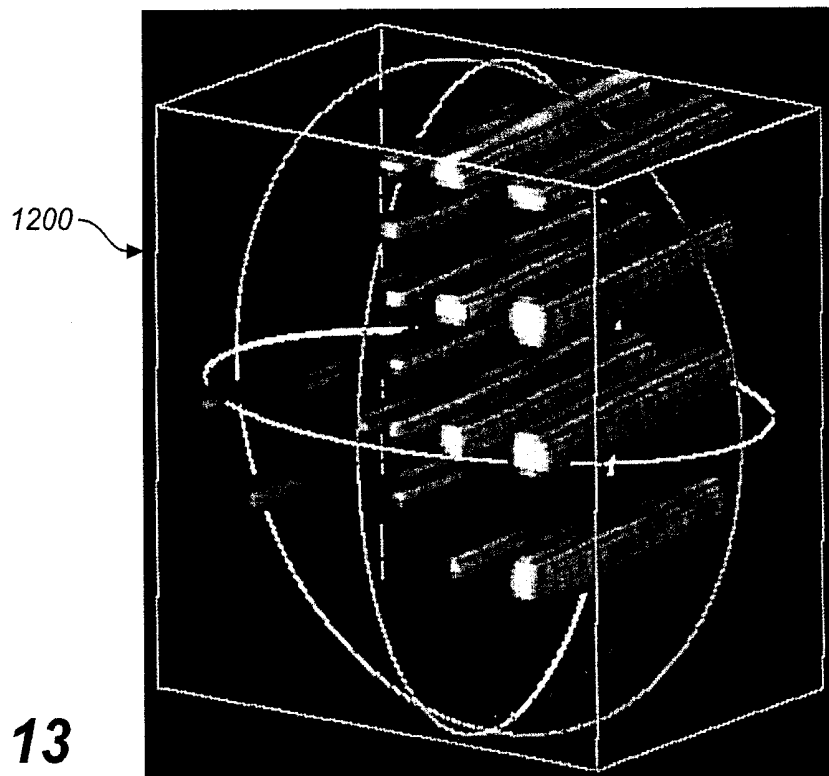


FIG. 12E

**FIG. 13**



# METHOD AND APPARATUS TO PRODUCE ULTRASONIC IMAGES USING MULTIPLE APERTURES

## CROSS REFERENCE TO RELATED APPLICATIONS

This application is a continuation of U.S. patent application Ser. No. 13/632,929, filed Oct. 1, 2012; which application is a continuation of U.S. patent application Ser. No. 13/215,966, filed Aug. 23, 2011, now U.S. Pat. No. 8,277,383; which application is a continuation of U.S. patent application Ser. No. 11/865,501, filed Oct. 1, 2007, now U.S. Pat. No. 8,007,439; which application claims the benefit of U.S. Provisional Patent Applications No. 60/862,951, filed Oct. 25, 2006, and U.S. Provisional Patent Applications No. 60/940,261, filed May 25, 2007; all of which are incorporated by reference in their entirety herein.

## BACKGROUND OF THE INVENTION

### 1. Field of the Invention

The present invention relates generally to imaging techniques used in medicine, and more particularly to medical ultrasound, and still more particularly to an apparatus for producing ultrasonic images using multiple apertures.

2. Discussion of Related Art Including Information Disclosed Under 37 CFR §§1.97, 1.98

In conventional ultrasonic imaging, a focused beam of ultrasound energy is transmitted into body tissues to be examined and the returned echoes are detected and plotted to form an image. In echocardiography the beam is usually stepped in increments of angle from a center probe position, and the echoes are plotted along lines representing the paths of the transmitted beams. In abdominal ultrasonography the beam is usually stepped laterally, generating parallel beam paths, and the returned echoes are plotted along parallel lines representing these paths. The following description will relate to the angular scanning technique for echocardiography (commonly referred to as a sector scan). However, the same concept with modifications can be implemented in abdominal scanners.

The basic principles of conventional ultrasonic imaging are well described in the first chapter of Echocardiography, by Harvey Feigenbaum (Lippincott Williams & Wilkins, 5<sup>th</sup> ed., Philadelphia, 1993). These will not be repeated here except as necessary to illustrate the differences between the conventional techniques and the present invention.

It is well known that the average velocity  $v$  of ultrasound in human tissue is about 1540 m/sec, the range in soft tissue being 1440 to 1670 m/sec (see for example P. N. T. Wells, Biomedical Ultrasonics, Academic Press, London, New York, San Francisco, 1977). Therefore, the depth of an impedance discontinuity generating an echo can be estimated as the round-trip time for the echo multiplied by  $v/2$ , and the amplitude is plotted at that depth along a line representing the path of the beam. After this has been done for all echoes along all beam paths, an image is formed, such as the image 10 shown in FIG. 1, in which a circle has been imaged. The gaps between the scan lines are typically filled in by interpolation. One of the earliest interpolation algorithms applied to echocardiography was described in U.S. Pat. No. 4,271,842, to Specht et al.

In order to insonify the body tissues, a beam formed either by a phased array or a shaped transducer is scanned over the tissues to be examined. Traditionally, the same transducer or array is used to detect the returning echoes. This design con-

figuration lies at the heart of one of the most significant limitations in the use of ultrasonic imaging for medical purposes; namely, poor lateral resolution. Theoretically the lateral resolution could be improved by increasing the aperture of the ultrasonic probe, but the practical problems involved with aperture size increase have kept apertures small and lateral resolution large. Unquestionably, ultrasonic imaging has been very useful even with this limitation, but it could be more effective with better resolution.

In the practice of cardiology, for example, the limitation on single aperture size is dictated by the space between the ribs (the intercostal spaces). For scanners intended for abdominal and other use, the limitation on aperture size is not so obvious, but it is a serious limitation nevertheless. The problem is that it is difficult to keep the elements of a large aperture array in phase because the speed of ultrasound transmission varies with the type of tissue between the probe and the area of interest. According to the book by Wells (cited above), the speed varies up to plus or minus 10% within the soft tissues. When the aperture is kept small, the intervening tissue is, to a first order of approximation, all the same and any variation is ignored. When the size of the aperture is increased to improve the lateral resolution, the additional elements of a phased array may be out of phase and may actually degrade the image rather than improving it. The instant disclosure teaches methods to maintain all of the information from an extended phased array "in phase" and thus to achieve sought-after improved lateral resolution.

In the case of cardiology, it has long been thought that extending the phased array into a second or third intercostal space would improve the lateral resolution, but this idea has met with two problems. First, elements over the ribs have to be eliminated, leaving a sparsely filled array. New theory is necessary to steer the beam emanating from such an array. Second, the tissue speed variation described above, but not adequately addressed until this time, needs to be compensated. The same solution taught in this disclosure is equally applicable for multi-aperture cardiac scanning, or for extended sparsely populated apertures for scans on other parts of the body.

## BRIEF SUMMARY OF THE INVENTION

The present invention solves both the problem of using more than one intercostal space and the problem of accommodating unknown phase delays from using elements spread over a large sparse aperture. The solution involves separating the insonifying probe from the imaging elements. The separation can be a physical separation or simply a separation in concept wherein some of the elements of the array can be shared for the two functions.

A single omni-directional receive element (such as a receive transducer) can gather all of the information necessary to reproduce a two-dimensional section of the body. Each time a pulse of ultrasound energy is transmitted along a particular path, the signal received by the omnidirectional probe can be recorded into a line of memory. [The terms "omni-directional probe," "omni probe" and/or "omni," are used synonymously herein to mean an omnidirectional probe.] When this is done for all of the lines in a sector scan, the memory can be used to reconstruct the image. This can be accomplished in the same time as data is being collected for the next frame.

There are numerous advantages to this approach, and these comprise the objects and advantages of the present invention. They include, among others:

The dominance of specular reflections so prominent in reconstructing images by returns to the main probe is greatly attenuated.

More than one omnidirectional probe can be used. Each one can be used to reconstruct an entire sector image but with different point spread functions. These can be combined to produce an image with a sharper point spread function.

Compensations can be made for different delays in different paths through the tissue.

Many more scan lines can be reconstructed than the number of pulses generated by the main probe. This overcomes the traditional limit of the number of scan lines by the speed of ultrasound in tissue, tissue depth of interest, and the time allowed between frames, which is typically  $\frac{1}{30}$ th second.

Artificial scan lines can be considered as overlapping, and each pixel on an output image can be imaged from information from more than one omni line of data. Therefore the output pixel can be averaged from multiple data, thus improving the signal-to-noise ratio.

Omnidirectional probes can be placed in multiple intercostal spaces, the suprasternal notch, the substernal window, multiple apertures along the abdomen and other parts of the body, and even on the end of a catheter. An advantage in using omnidirectional probes for catheter placement is that no steering is required of the probe.

Probes can be placed either on the image plane, off of it, or any combination. When placed away from the image plane, omni probe information can be used to narrow the thickness of the sector scanned.

There has thus been broadly outlined the more important features of the invention in order that the detailed description that follows may be better understood, and in order that the present contribution to the art may be better appreciated. Additional objects, advantages and novel features of the invention will be set forth in part in the description as follows, and in part will become apparent to those skilled in the art upon examination of the following. Furthermore, such objects, advantages and features may be learned by practice of the invention, or may be realized and attained by means of the instrumentalities and combinations particularly pointed out in the appended claims.

Still other objects and advantages of the present invention will become readily apparent to those skilled in this art from the following detailed description, which shows and describes only the preferred embodiments of the invention, simply by way of illustration of the best mode now contemplated of carrying out the invention. As will be realized, the invention is capable of modification in various obvious respects without departing from the invention. Accordingly, the drawings and description of the preferred embodiment are to be regarded as illustrative in nature, and not as restrictive.

#### BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWINGS

FIG. 1 is a diagrammatic view of a simulation showing a circular object imaged by a conventional sector scanner.

FIG. 2 is a schematic diagram of the axes representing the relative positions of the insonifying and omni-directional probes.

FIG. 3 is a graph showing the orientation of the point spread function of an imaging system using a single omnidirectional probe as a function of the position of the omnidirectional probe.

FIG. 4 is a graph showing the orientation of the point spread function of an imaging system using a single omnidi-

rectional probe as a function of the position of the omnidirectional probe when it is not in the scan plane.

FIG. 5 is a diagrammatic view of a simulation showing the same circular object of FIG. 1 imaged by data received by a single omni element located at  $x_0=40$  mm,  $y_0=0$  mm,  $z_0=0$  mm.

FIG. 6 is a schematic diagram illustrating the relative positions of probes showing, in addition, points A-A', which have equal round trip distances and times.

FIG. 7 is a schematic diagram showing a possible fixture for positioning an omni-directional probe relative to the main (insonifying) probe.

FIG. 8 is a schematic diagram showing a non-instrumented linkage for two probes.

FIG. 9 is a schematic diagram showing variables for computation of x and y positions from received echoes.

FIG. 10A is a phantom image taken with a standard Acuson 128 XP-10 with a 3.5 MHz transducer and harmonic processing.

FIG. 10B is the same phantom image as that shown in FIG. 10A and taken with the same XP-10, wherein the center 64 elements were obscured but external processing employed to show improved lateral resolution. The progressions of anechoic areas on the phantom are 8 mm diameter, 6 mm, 4 mm, 3 mm, and 2 mm.

FIG. 11 is an image of the same phantom produced by the same transducer as the images in FIGS. 10A and 10B, with the center obscured, but with substantial image processing over multiple scans. Note that even though the total aperture is only 19 mm that the 2 mm diameter anechoic areas are now visible. Lateral resolution could be greatly improved if the two parts of the transducer were physically separated and the phased delays reprogrammed for the resulting geometry.

FIG. 12A is a schematic perspective view showing an adjustable, extendable hand held two-aperture probe (especially adapted for use in cardiology US imaging). This view shows the probe in a partially extended configuration.

FIG. 12B is a side view in elevation thereof showing the probe in a collapsed configuration.

FIG. 12C shows the probe extended so as to place the heads at a maximum separation distance permitted under the probe design, and poised for pushing the separated probe apertures into a collapsed configuration.

FIG. 12D is a side view in elevation again showing the probe in a collapsed configuration, with adjustment means shown (i.e., as scroll wheel).

FIG. 12E is a detailed perspective view showing surface features at the gripping portion of the probe.

FIG. 13 is a 3D image highlighting the anechoic tubes of the ATS Model 539 phantom.

#### DETAILED DESCRIPTION OF THE INVENTION

A key element of the present invention is that returned echoes in ultrasonography can be detected by a separate relatively non-directional receive transducer located away from the insonifying probe (transmit transducer), and the non-directional receive transducer can be placed in a different acoustic window from the insonifying probe. This probe will be called an omni-directional probe because it can be designed to be sensitive to a wide field of view.

If the echoes detected at the omni probe are stored separately for every pulse from the insonifying transducer, it is surprising to note that the entire two-dimensional image can be formed from the information received by the one omni.

Additional copies of the image can be formed by additional omni-directional probes collecting data from the same set of insonifying pulses.

A large amount of straightforward computation is required to plot the amplitude of echoes received from the omni. Referring now to FIG. 2, in which there is shown the position of the omni-directional probe **100** relative to the position of the insonifying (main) probe **120** and the insonifying beam **130**. The position of the omni-directional probe relative to the beam is indicated by  $x_0$ ,  $y_0$  and  $z_0$  **140**, where  $x_0$  and  $y_0$  are in the scan plane **150** scanned by the insonifying beam and  $z_0$  is distance perpendicular to that plane. Instead of simply plotting the depth along the scan line  $d=tv/2$  (where  $t$  is the round-trip time, it is now computed as that point  $d=\sqrt{x^2+y^2}$  for which  $tv=d+\sqrt{(x-x_0)^2+(y-y_0)^2+z_0^2}$ ).

This procedure will produce a sector scan image similar to that using the conventional technique except that the point spread function will be rotated. FIG. 3 shows the orientation **200** of the psf as a function of the position of the omni probe. A single point at  $x=0$ ,  $y=70$  mm,  $z=0$  is the point being imaged, and the groups of dots **210** each indicate the orientation of the psf if the omni-probe were placed at the location of the center of each group. The insonifying probe (main probe) is located at the center group **220** on the bottom of the figure. In this simulation the horizontal (x axis) **230** shown goes from -40 mm to +40 mm. The vertical (y axis) **240** goes from 0 to 80 mm depth.

FIG. 4 shows the orientation **300** of the psf as a function of the position of the omni probe if it is not in the scan plane. In this simulation the horizontal (x axis) **310** shown again goes from -40 mm to +40 mm, but the vertical axis **320** is the z axis (distance away from the scan plane) and goes from 0 to 80 mm away from the scan plane.

FIG. 5 shows a plot of the same circular object **20** as in FIG. 1, plotted, however, from data received by a single omni element. Other complete two dimensional reconstructions may be formed using data from additional omni elements, if desired.

Specular reflection is reduced using the omni probe compared to using the main probe for both insonification and detection. This is because all parts of a surface normal to the main beam are insonified with the same phase. When the phase is such that maximum echo is returned, all the echoes add to produce a specular echo. When the signals are normalized to accommodate the dynamic response of a particular display device, non-specular echoes will tend to drop out.

In contrast to this, and referring now to FIG. 6, there is also shown the relative positions of probes, but there is also shown points A-A' **400**, which have equal round trip distances and times. Points A-A' are on a surface normal to the bisector line  $a_1=a_2$  **410**, which also have equal round trip distances and times, are not insonified with the same phase, and do not all reflect equally. This attenuation of the specular reflection is particularly important when visualizing circular structures which frequently have surfaces normal to the main beam.

An algorithm to plot this data on a rectangular grid is: (a) for each point on the x,y grid, convert x,y to depth and angle; (b) then find closest angle  $k$  scanned by the insonifying beam; (c) if it is sufficiently close, then convert x,y to distance to the omni; (d) compute time  $t=(\text{distance to insonifying beam} + \text{distance to omni})/v$ ; and (e) plot amplitude recorded by the omni for the  $k$  scan at x,y.

However, more information is available and should be used. It is possible to use the same technique to plot additional scan lines which were not explicitly insonified by the insonifying probe. Because of the inherently wide beam width of medical ultrasonic probes, much tissue between intentional

scan lines is also insonified and returns echoes. Making use of this information is particularly important when capturing motion (especially in echocardiography) because the number of pulses that can be generated is strictly limited by the speed of ultrasound in tissue and the scan repetition rate desired.

The reconstructed image will get better as the angle between the main beam and the omni gets larger. However it is not necessary to focus a narrow beam on every element of tissue to be imaged as is true if the data is not stored and then processed before display. The lateral resolution can be reconstructed using a Wiener filter to be much better than the beam width if the noise spectrum is low enough. In one simulation of 2 circles of diameter 2.2 mm and 4.0 mm, both imaged well enough that the center was clear even though the beam width was 4.4 mm tapered from 1 to 0 by a cosine function. The Wiener filter is described in the next section.

There are four main sources of noise in ultrasonic imaging: (1) blur due to array size not wide enough; (2) shot noise; (3) reverberation from big interfaces; and (4) speckle.

Multiple probes give independent measures of shot noise, but using closely spaced elements in the main probe (if it is a phased array) will not give independent noise for the other three sources. Adding one or more omni probes will change the look angle, which will thereby change the speckle pattern and the reverberation pattern. These can be averaged out to lower the noise power spectrum. The Wiener filter can then be employed to cancel the blur.

Another way to eliminate speckle is to obtain a good sample of it for estimates of the noise spectrum to then be used in the Wiener filter.

De-blurring and de-noising by these techniques using only an external omni probe or probes will make it possible to visualize small and moving objects such as the coronary arteries. In such a case medical personnel could assess the degree of opening in the lumen or patency of bypass grafts without resort to invasive catheterization techniques.

When combining more than one image such as one from the main probe and another from one or more omni probes for the purpose of averaging out the various sources of noise, it is necessary to compensate for the variation in ultrasound velocity through different paths. Experiments have shown that small unaccounted errors in path velocity will displace the reconstructed image in both horizontal and vertical positions. Cross correlation techniques should be used to find the displacement with one image taken as reference before addition or other combination of images.

Two possibilities exist with regard to the Wiener filter. In one, a Wiener filter can be used separately on each image and then combine them. Alternatively, one can first combine the images (yielding a more complex point spread function) and then employ Wiener filtering.

In order to perform the indicated computations, it is necessary to either measure or estimate the position ( $x_0$ ,  $y_0$ ,  $z_0$ ) of the omni relative to the main probe. Exact knowledge of these positions is not required because, as we have seen, the displacement of the image from the omni probe is also affected by the variation of the velocity of ultrasound in different types of tissue. For this reason it is necessary to use cross correlation or some other matching criterion to make a final correction of the position of the omni-generated image before combining with the reference image or images.

Determining the Position of the Omni Probe(s): There are many ways to determine the position of the omni probe. Referring now to FIG. 7, one way is to provide apparatus **500** for pivotally and/or swivelingly mounting the omni probe **510** on a fixture **520** attached to the insonifying probe **530**. The fixture preferably includes articulated joints **540** with sensors

(not shown) to measure angles and distances **550** of the links. FIG. 7 illustrates a simplified version of such a fixture, wherein fixed hinges allow movement of only  $x_0$  and  $y_0$ .

Another method is to have no mechanical connection between the omni probe and the main probe (except wires for signals and power). Instead, the omni probe can transmit a signal using radio frequencies, light, or a different frequency ultrasound to triangulation receivers mounted on the main probe or a separate platform.

A third method again has no mechanical connection between the omni probe and main probe. For this method the omni probe (or probes) can be attached to the patient with tape, and the ultrasonographer can manipulate the main probe to find the best image without regard to positioning the omni probe(s). As indicated in FIG. 4, a two-dimensional image can be formed separately from the echoes received from the omni probe and from the main probe. By adjusting four variables ( $x_0$ ,  $y_0$ ,  $z_0$  and D, the average difference in time for ultrasound to go through different tissue types instead of traveling through idealized tissue of constant ultrasound velocity), the images can be made to coincide. The four variables can be adjusted iteratively to maximize cross-correlation or another measure of similarity. Standard multi-dimensional search techniques that could be used include gradient ascent, second order (Newton's method), simulated annealing, and genetic algorithms. A fifth variable, the angle of the psf which is necessary for deconvolution, is implied from the first four variables. Misregistration of the images can be caused by inaccurate estimation of any of the four variables, but good registration can be achieved by simply adjusting D which will tend to compensate for errors in estimates of the others.

When the application requires the highest resolution compatible with capturing motion at a high frame rate, the four variables can be estimated over several frames of information. When the ultrasonographer has selected a good view angle, the frames can be combined at high rate holding  $x_0$ ,  $y_0$ ,  $z_0$  and D constant.

When the application requires the highest possible resolution, data can be captured (perhaps with EKG gating to capture separate images at systole and diastole) and the multi-dimensional search to optimize matching can be done more accurately although not in real time. Two advantages of this approach is that different values of D can be found for systole and diastole, and that different psf's can be used for deconvolution at different depths in the image.

Determining the Position of the Omni Probe(s) Using Correlation of the Scan Line Data Rather than Complete 2D Sectors: A fourth method for determining the position of the omni probe(s) entails replacing the omni probe or probes with a "semi-omni probe" or probes. The reason for this is to increase the signal to noise ratio by restricting the sensitive region of the receive transducer to a plane rather than a hemisphere. Because of this restriction it is necessary to have a mechanical linkage to ensure that both the transmit and receive transducers are focused on the same plane.

Two probes could be placed in any two acoustic windows. In the case of echocardiography, one would likely be in the normal parasternal window which typically gives the clearest view of the whole heart. A second window available in most patients is the apical view. Another window usually available is the subcostal. The two probes could even be placed on either side of the sternum or in parasternal windows in two intercostal spaces.

One probe could be the standard phased array cardiac probe. The second (and third, etc.) would be used as receive only. Theoretically it could be omnidirectional, but that

would necessarily provide lower signals and therefore low signal to noise ratios (S/N). A better alternative is to use a probe which is ground to be sensitive to a plane of scan but omnidirectional within that plane. A single piece of PZT would work well, but to minimize the amount of new design required it is also possible to use a second probe head similar to the main probe and then use individual elements or small groups combined to act as single elements. The design goal is to use as many elements as possible to maximize signal to noise ratio while using few enough to minimize angle sensitivity.

In this embodiment **600** (see FIG. 8), the two probes **610**, **620**, may be linked together with an articulating mechanical linkage, which ensures that the plane of scan of each probe includes the other, but the distance between them is unconstrained. A slave servomechanism is also possible, but the mechanical linkage will be described here.

The procedure is to aim the main probe **620** at the target **630** (e.g., heart) and position the secondary probe **610** at a second window with maximum received signal strength. One possibility is that the main probe be positioned for a long axis view with the secondary probe over the apex of the heart. Some slight deviation of the long axis view may be necessary in order to maintain the secondary probe in its most sensitive spot.

The secondary probe would now be held on the patient with a mechanical housing which allows a fan or rocking motion. The disadvantage of having two probes in fixed positions on the body is that the plane of scan must include these two points. The only degree of freedom is the angle at which the scan plane enters the patient's body. For a conventional 2D examination this is a severe limitation, but if the goal is to gather three-dimensional information, this is not a limitation. The 3D information is obtained by rocking the main probe back and forth through a sufficient angle so that the entire heart isinsonified. The secondary probe also rocks back and forth by virtue of the mechanical linkage between the probes. The instantaneous angle of rocking must be monitored—perhaps by reference to a gyroscope mounted with the main probe. The rocking could be actuated by the hand motion of the ultrasound technician, or it could be motorized for a more-uniform angle rate. In an alternative preferred embodiment (for echocardiography), the main probe and an array of omni probes are placed in adjacent intercostal spaces using a mechanism as shown in FIG. 12.

Computer software could be provided such that the 2D slices would fill a 3D volume of voxels. After adjacent voxels are filled through interpolation, 3D information can be displayed as projections or as slices through the volume at arbitrary orientations.

The need for and one important use of the 3D information is covered in U.S. patent application Ser. No. 11/532,013, now U.S. Pat. No. 8,105,239, also by the present inventor, and which application is incorporated in its entirety by reference herein.

Yet another variation on this theme is to have the secondary transducer mechanically linked to the primary so that each plane of scan contains the other transducer (as above), but allow rotation of the main probe about its own axis. In this case the secondary probe would be allowed to move on the patient's body (properly prepared with ultrasound gel). It would have many elements, and an attached computer would scan them all to find those elements which have the strongest return signal.

Estimating Relative Probe Positions from Reflected Signals: For image reconstruction it is essential to know the position of the secondary probe ( $x$ ,  $y$ ) relative to the main

probe. This has to be evaluated separately for each frame of data because of the motion of the patient, technician, and/or motorized angle actuator. Since the linkage will prevent any difference in position (z) perpendicular to the scan plane, only x and y need be assessed.

Note that any tilt of the main probe will change the reference axes so that x and particularly y will change too.

When a pulse is transmitted from the main probe it insonifies a sequence of tissues in the path of the beam. The returns from the tissues will be received by both the main probe and the secondary, digitized and stored in the computer. Echoes from relatively proximate tissues will be different for the two probes, but echoes from mid- to far range will be similar. It is possible to use cross correlation to find similar small patches in the two stored returns. They will be similar except for the time delay relative to the launching of the pulse from the main probe. The time delay will be related to the offsets x and y. Values for x and y cannot be determined from one set of time delays, but can be determined by solving a set of simultaneous equations from two detected similar returns. These could be different patches of the same pulse return or from returns from differently directed main pulses.

Referring now to FIG. 9, if the main probe 700 transmits at angle  $90^\circ - \alpha$  relative to its centerline and an identifiable packet of returns occurs at time  $t_{1m}$  at the main probe and at time  $t_{1s}$  at the secondary (omni) probe 720, then:

tissue packet at  $(x_1, y_1)$  is received at time  $t_{1m}$ , and distance  $m_1 = \sqrt{x_1^2 + y_1^2}$

tissue packet at  $(x_2, y_2)$  is received at time  $t_{2m}$ , and distance  $m_2 = \sqrt{x_2^2 + y_2^2}$

$t_{1m}$  corresponds to time of two trips of distance  $m_1$

$t_{1m}s = 2m_1$ , where  $s$  = speed of ultrasound in same units as  $m$  = approx.  $1.54 \times 10^6$  mm/sec

$t_{1s}s = m_1 + \sqrt{(x_1 - x)^2 + (y_1 - y)^2}$ .

Similarly,  $t_{2m}s = 2m_2$

$$t_{2s}s = m_2 + \sqrt{(x_2 - x)^2 + (y_2 - y)^2}$$

$$(t_{1s}s - 0.5t_{1m}s)^2 = (x_1 - x)^2 + (y_1 - y)^2$$

$$(t_{2s}s - 0.5t_{2m}s)^2 = (x_2 - x)^2 + (y_2 - y)^2 \quad (1)$$

Since  $X_1, y_1, x_2, y_2$  and the times are known, one can solve the last two simultaneous equations for x and y. Similarly, if a z offset between the two probes is allowed, x, y, and z can be calculated by solving three simultaneous equations.

Many more measurements from packet pairs are available. One could make a measurement on several or every scan line (angle) as measured from the main probe. Then we would have many equations in 2 unknowns which can be used to make more-accurate estimations of the 2 unknowns. Since these are nonlinear equations, a search technique can be utilized. One way to accomplish this is to compute error squared over a grid of (x, y) points using the equation:

$$E_2 = \sum_{i=1}^N \left( \sqrt{(x_i - x)^2 + (y_i - y)^2} - t_{is} + 0.5t_{im}s \right)^2 \quad (2)$$

The minimum  $E_2$  will indicate the minimum squared error estimate of (x, y). The search should be conducted over the expected range of x and y to save time and to avoid spurious ambiguous minima.

When the z component of the relative position is not constrained to be zero, the comparable error squared equation is:

$$E_2 = \sum_{i=1}^N \left( \sqrt{(x_i - x)^2 + (y_i - y)^2 + z_i^2} - t_{is} + 0.5t_{im}s \right)^2$$

The minimum  $E_2$  will indicate the minimum squared error estimate of (x, y, z).

If the speed of sound on the return path to the secondary (omni) transducer is different from  $s$  due to different types of tissues being traversed, the values of x and y (and z if used) will be different from the geometric values. However, use of these values in the image reconstruction algorithm will automatically compensate for the different speeds.

Obviously, the probes that have been described for imaging the heart would work equally well for imaging abdominal organs and other parts of the body such as legs, arms, and neck. In fact, use of receive-only transducers in conjunction with a transmit/receive probe would work better for abdominal organs because the orientation of the probe set is not limited by the intercostal spaces formed by the ribs. Whereas the locations of the acoustic windows to the heart limit the orientation of the probe to only a few orientations and it is necessary to rock the probe to gather three dimensional data, the probes can be used on the abdomen in any orientation presently used. Therefore the probes can be used for real-time 2D scans to duplicate presently accepted procedures except with much higher lateral resolution. In fact, this application of the technology may be as important as the application to cardiology (which was our original motivation).

For abdominal scanning it is not necessary to have an elaborate spacing adjustment between the active transmit/receive elements and the receive-only elements. In fact they could all be mounted together in one rigid probe, either as a linear array or an array with known curvature. Some prior art wide linear arrays exist which insonify tissue by using a small subset of the total number of elements to transmit and receive a beam perpendicular to the array. Then another partially overlapping subset of elements is used to transmit and receive another line parallel to the first one, and so on until an entire scan is completed.

However, the same array could be partitioned into an active section plus one or more passive sections where all sections would be used for each pulse. The active section of elements would be used in transmit as a sector scanner sending out beams in a sequence of angular paths. On receive, all elements would be treated as independent relatively nondirectional receivers and their outputs would be combined to form a high resolution image by the methods taught in this patent. Cross-correlation image matching to account for the variations in ultrasound speeds could be done separately for each receive element or for groups of elements for which the speed corrections would be nearly the same.

The concept of mounting the active and receive-only elements on a rigid structure eliminates the necessity for articulating and instrumenting the spacing between elements thus making practical combined probes to be used for trans-esophageal (TEE), trans-vaginal, and trans-rectal imaging.

A final class of probes would involve putting a receive-only transducer or transducers on the end of a catheter to be inserted in an artery, vein, or urethra while a separated transmit transducer array is applied to the surface of the skin. The advantage of this approach is that the catheter could be positioned close to an organ of interest thereby reducing the total transit distance from the transmit transducer to the receive element and thus higher frequencies could be used for better

resolution. The receive element(s) on the catheter would not have to be steered as it (they) would be relatively omnidirectional.

The Wiener Filter: The Wiener filter itself is not new, but since it is important for the de-convolution step it will be described briefly here in the context of the present invention. The Wiener filter is the mean squared error optimal stationary linear filter for images degraded by additive noise and blurring. Wiener filters are usually applied in the frequency domain. Given a degraded image  $I(u,v)$ , one takes the discrete Fourier Transform (DFT) or the Fast Fourier Transform (FFT) to obtain  $I(u,v)$ . The true image spectrum is estimated by taking the product of  $I(u,v)$  with the Wiener filter  $G(u,v)$ :

$$\hat{S} = G(u,v)I(u,v)$$

The inverse DFT or FFT is then used to obtain the image estimate  $s(n,m)$  from its spectrum. The Wiener filter is defined in terms of the following spectra:

(a)  $H(u,v)$ —Fourier transform of the point spread function (psf);

(b)  $P_s(u,v)$ —Power spectrum of the signal process, obtained by taking the Fourier transform of the signal autocorrelation;

(c)  $P_n(u,v)$ —Power spectrum of the noise process, obtained by taking the Fourier transform of the noise autocorrelation;

The Wiener filter is:

$$G(u,v) = \frac{H^*(u,v)P_s(u,v)}{|H(u,v)|^2P_s(u,v) + P_n(u,v)}$$

The ratio  $P_s/P_n$  can be interpreted as signal-to-noise ratio. At frequencies with high signal to noise ratio, the Wiener filter becomes  $H^{-1}(u,v)$ , the inverse filter for the psf. At frequencies for which the signal to noise ratio is low, the Wiener filter tends to 0 and blocks them out.

$P_s(u,v) + P_n(u,v) = |I(u,v)|^2$ . The right hand function is easy to compute from the Fourier transform of the observed data.  $P_n(u,v)$  is often assumed to be constant over  $(u,v)$ . It is then subtracted from the total to yield  $P_s(u,v)$ .

The psf can be measured by observing a wire phantom in a tank using the ultrasound instrument. The Fourier transform of the psf can then be stored for later use in the Wiener filter when examining patients.

Because the psf is not constant as a function of range, the Wiener filter will have to be applied separately for several range zones and the resulting images will have to be pieced together to form one image for display. A useful compromise might be to optimize the Wiener filter just for the range of the object of interest such as a coronary artery or valve. It will be necessary to store separate Wiener filters for each omnidirectional probe and for the main probe when it is used as a receive transducer.

An alternative to the Wiener Filter for deconvolution is the least mean square (LMS) adaptive filter described in U.S. patent application Ser. No. 11/532,013, now U.S. Pat. No. 8,105,239. LMS Filtering is used in the spatial domain rather than the frequency domain, and can be applied to the radial scan line data, the lateral data at each depth, or both together.

Image sharpening can be accomplished by the use of unsharp masking. Because aperture blur is much more pronounced in the lateral dimension perpendicular to the insonifying beam) than in the radial dimension, it is necessary to perform unsharp masking in only one dimension. When using a sector scanner, this masking should be performed before

scan conversion. When using a linear phased array, the unsharp masking should be performed on each data set of constant range. Unsharp masking consists of intentionally blurring an image, subtracting the result from the original image, multiplying the difference by an arbitrary factor, and adding this to the original image. In one dimension this is the same as blurring a line of data, subtracting it from the original line, and adding a multiple of the difference to the original line.

Multiple Active Transducers—Two Alternative Approaches: It is possible to use more than one active transducer placed at multiple acoustic windows in order to achieve the same goals of increased lateral resolution and noise suppression. A practical method of providing multiple omni probes is to use a second phased array head in a second acoustic window and then treating each element or group of elements of the second phased array as a separate omni. With this configuration of probes it would be possible to switch the functions of the two probe heads on alternate scans thereby generating images with different speckle patterns which can be averaged out.

Multiple phased array heads can also be used together so that both are active on the same scan. When two (or more) phased array transducers are placed in the same scan plane, they can be programmed with delays such that they act as a single array with a gap in the array of transducer elements. The advantages of having a gap in the array include a) achieving the lateral resolution of a wide aperture without the expense of filling in acoustic elements through the gap, and b) the gap in the probe or between probes can be fitted over ribs or the sternum. The first advantage applies equally to applications other than cardiac. The disadvantages of multiple active probes is that both the transmit and receive delays have to be recomputed for each new gap dimension and/or angular orientation of one probe relative to the others.

An active probe with a gap has been demonstrated to produce lateral resolution as good as the probe without the gap. This implies that larger gaps will achieve higher resolution since lateral resolution is determined primarily by the overall aperture. Referring now to FIG. 10A, there is illustrated the image 800 of an ATS Laboratories Model 539 phantom was imaged using an Acuson 128 XP-10 ultrasonic scanner with a 4V2c probe. In FIG. 10B, the same probe was used to image the phantom with its center 64 elements totally obscured by aluminum foil and electrical tape. As can be seen, the lateral resolution in the image 900 is as good as the original although the image quality is degraded by speckle and other noise.

FIG. 11 shows an image 1000 of the same phantom produced by the same transducer with the center obscured, but with substantial image processing over multiple scans.

FIGS. 12A-E are various views showing an adjustable, extendable hand held two-aperture probe 1100 adapted for use in cardiology US imaging. This apparatus embodies the inventive concept of separating the insonifying probe 1110 (a transmit transducer) from the imaging elements 1120 (receiver transducer). This comfortable device includes adjustment means 1130, such as a scroll wheel, which selectively drives the elements either closer or further apart along either a medial telescoping portion 1140 or a medial insertable sleeve, and thereby provides a range of separation at predetermined distances. The gripping portion 1150 provides easy access to the scroll wheel and places the user's hand in the functional position to minimize overuse injury. FIG. 12A shows the probe in a partially extended configuration. FIG. 12B shows the probe in a collapsed configuration. FIG. 12C shows the probe extended so as to place the heads at a maximum separation distance permitted under the probe design.

13

FIG. 12D shows the probe in a collapsed configuration, with adjustment means shown. And FIG. 12E is a detailed perspective view showing surface features at the gripping portion of the probe.

FIG. 13 shows a three-dimensional display 1200 of the anechoic tubes of the Model 539 phantom. This 3D display was formed from 13 parallel slices produced with the same transducer with the center obscured. When the total aperture is increased it will be possible to display smaller anechoic tubes such as the coronary arteries. The processing involved for this display is a combination of the techniques of the instant patent and those of U.S. patent application Ser. No. 11/532,013 now U.S. Pat. No. 8,105,239.

Having fully described several embodiments of the present invention, many other equivalents and alternative embodiments will be apparent to those skilled in the art. These and other equivalents and alternatives are intended to be included within the scope of the present invention.

What is claimed is:

1. A method of ultrasound imaging, comprising:
  - transmitting a first ultrasound signal from a transmit transducer array into a target object;
  - receiving first echoes of the first ultrasound signal with a first transducer that is omnidirectional in an image plane and is located in a first acoustic window;
  - storing the first echoes in a computer memory;
  - receiving second echoes of the first ultrasound signal with a second transducer that is omnidirectional in the image plane and is located in a second acoustic window that does not overlap the first acoustic window;
  - storing the second echoes in the computer memory;
  - obtaining position information of the first and second transducers relative to the transmit transducer array;

14

retrieving the first echoes from the computer memory and constructing a first image of the target object from the retrieved first echoes and the position information of the first transducer;

retrieving the second echoes from the computer memory and constructing a second image of the target object from the retrieved second echoes and the position information of the second transducer; and  
combining the first and second images to form a combined image.

2. The method of claim 1, further comprising, prior to combining the first and second images, determining a displacement of the second image relative to the first image.

3. The method of claim 1, further comprising applying a Wiener filter to the first and second images before combining the first and second images.

4. The method of claim 1, further comprising applying a Wiener filter to the combined image.

5. The method of claim 1, wherein the second transducer is at least one transducer element on the transmit transducer.

6. The method of claim 1, wherein the second transducer is separate from the transmit transducer and the first transducer.

7. The method of claim 1, wherein the target object is human tissue.

8. The method of claim 7, wherein the first acoustic window is selected from the group consisting of a first parasternal intercostal space, a second parasternal intercostal space, a suprasternal notch, a substernal position, a subcostal window, an apical view, a first intercostal space adjacent the sternum, and a second intercostal space adjacent the sternum, and wherein the second acoustic window is selected from the same group.

\* \* \* \* \*

专利名称(译)	使用多个孔产生超声图像的方法和设备		
公开(公告)号	<a href="#">US9072495</a>	公开(公告)日	2015-07-07
申请号	US14/157257	申请日	2014-01-16
[标]申请(专利权)人(译)	SPECHT DONALD F		
申请(专利权)人(译)	SPECHT唐纳德F.		
当前申请(专利权)人(译)	MAUI IMAGING , INC.		
[标]发明人	SPECHT DONALD F		
发明人	SPECHT, DONALD F.		
IPC分类号	A61B8/14 A61B5/00 G01S15/89 A61B8/00 A61B8/08 G01S7/52		
CPC分类号	A61B8/5253 A61B8/42 A61B8/4209 A61B8/4281 A61B8/4455 A61B8/483 G01S7/52046 G01S15/8977 A61B5/725 A61B8/085 A61B8/145 A61B8/4494 A61B8/5207 A61B8/543 G01S7/5205 A61B8/4218 A61B8/4483 A61B8/08 A61B8/4245		
优先权	13/215966 2012-10-02 US 11/865501 2011-08-30 US 60/862951 2006-10-25 US 60/940261 2007-05-25 US		
其他公开文献	US20140135626A1		
外部链接	<a href="#">Espacenet</a> <a href="#">USPTO</a>		

#### 摘要(译)

描述了超声波扫描器和全向接收换能器的组合，用于从接收到的回波产生二维图像。具有不同噪声分量的二维图像可以由附加换能器接收的回波构成。这些可以组合以产生具有更好的信噪比和横向分辨率的图像。还公开了一种基于信息内容的方法，以补偿通过介入组织的不同路径的不同延迟。所公开的技术在医学成像中具有广泛的应用，但是理想地适合于使用两个或更多个肋间空间的多孔径心脏成像。由于横向分辨率主要由端部元件限定的孔径确定，因此不必用等间隔的元件填充整个孔径。使用这些方法的多个切片可以组合以形成三维图像。

