

OCT 27 2004

510(K) SUMMARY

G20 Diagnostic Ultrasound system

This summary of safety and effectiveness is provided as part of this Premarket Notification in compliance with the Safe Medical Device Act of 1990 revisions to 21 CFR, Part 807.92, Content and Format of a 510(k) Summary.

1. **Submitted By:**

Siemens Medical Solutions USA, Inc., Ultrasound Division  
22010 S.E. 51st Street  
Issaquah, WA 98029

**Contact Person:**

Patrick J Lynch  
Regulatory Affairs

Phone: (425) 557-1825

FAX: (425) 391-9198

**Date Prepared:**

September 20, 2004

2. **Proprietary Name:**

SONOLINE G20™ Diagnostic Ultrasound System

**Common/ Usual Name:**

Diagnostic Ultrasound System with Accessories

**Classification Name:**

21 CFR 892.1550

Ultrasonic Pulsed Doppler Imaging System      FR # 892.1550      Product Code 90-IYN

Ultrasonic Pulsed Echo Imaging System      FR # 892.1560      Product Code 90-IYO

Diagnostic Ultrasound Transducer      FR # 892.1570      Product Code 90-ITX

3. **Predicate Device:**

K040502, 03/09/2004, SONOLINE G20 Diagnostic Ultrasound System  
K020353, 02/13/2002, SONOLINE G50 & G60 S Diagnostic Ultrasound Systems  
K946179, 10/03/1995, marketed as SONOLINE Adara Diagnostic Ultrasound System

4. **Device Description:**

The G20 is a general purpose, mobile, software-controlled, diagnostic ultrasound system with an on-screen display for thermal and mechanical indices related to potential bioeffect mechanisms. Its function is to acquire primary or secondary harmonic ultrasound echo data and display it in: B-Mode, M-Mode, a combination of modes, 3D imaging or Harmonic Imaging on a CRT display.

The G20, has been designed to meet the following product safety standards:

- UL 60601-1, Safety Requirements for Medical Equipment
- CSA C22.2 No. 601-1, Safety Requirements for Medical Equipment
- AIUM/NEMA, 1998, Standard for Real Time Display of Thermal and Mechanical Acoustic Output Indices on Diagnostic Ultrasound Equipment
- AIUM/NEMA UD-2, 1998 Acoustic Output Measurement Standard for Diagnostic Ultrasound
- 93/42/EEC Medical Devices Directive
- Safety and EMC Requirements for Medical Equipment

- EN 60601-1
- EN 60601-1-1
- EN 60601-1-2
- EN 60601-2-37
- IEC 1157 Declaration of Acoustic Power
- ISO 10993 Biocompatibility

5. **Intended Uses:**

The G20 ultrasound imaging system is intended for the following applications: General Radiology, Abdominal, Small Parts, Transcranial, OB/GYN, Cardiac, Pelvic, Neonatal/Adult Cephalic, Urology, Vascular, Musculoskeletal, Superficial Musculoskeletal, and Peripheral Vascular applications.

The system also provides for the measurement of anatomical structures and for analysis packages that provide information that is used for clinical diagnosis purposes.

6. **Technological Comparison to Predicate Device:**

The G20 is substantially equivalent to the SONOLINE Adara, cleared via K946179, the SONOLINE G50/G60 S, cleared via K020353, and the SONOLINE G20, cleared via K040502. All systems transmit ultrasonic energy into patients, then perform post processing of received echoes to generate on-screen display of anatomic structures and fluid flow within the body. All systems allow for specialized measurements of structures and flow, and calculations.

**End of 510(k) Summary**



OCT 27 2004

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Siemens Medical Solutions USA, Inc.  
% Mr. Mark Job  
Responsible Third Party Official  
Regulatory Technology Services LLC  
1394 25<sup>th</sup> Street NW  
BUFFALO MN 55313

Re: K042833  
Trade Name: SONOLINE G20<sup>TM</sup> Diagnostic Ultrasound System  
Regulation Number: 21 CFR 892.1560  
Regulation Name: Ultrasonic pulsed echo imaging system  
Regulation Number: 21 CFR 892.1570  
Regulation Name: Diagnostic ultrasonic transducer  
Regulatory Class: II  
Product Code: 90 IYO and ITX  
Dated: October 11, 2004  
Received: October 13, 2004

Dear Mr. Job:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

This determination of substantial equivalence applies to the following transducers intended for use with the SONOLINE G20<sup>TM</sup> Diagnostic Ultrasound System, as described in your premarket notification:

Transducer Model Number

C5-2 Convex Array

L10-5 Linear Array

EV9-4 Convex Array

7.5L75S Linear Array

EC9-4 Convex Array Endocavity  
C4-2 Convex Array  
C8-5 Convex Array  
BE9-4 Biplane Endocavity  
C6F3 Convex Array Mechanically Driven, 3D  
EV8F5 Mechanical Sector Endovaginal 3D  
3.5C40S Convex Array  
5.0C40S Convex Array  
Endo PII Mechanical Sector Biplane Endo-cavity  
Endo VII Mechanical Sector Endovaginal

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This determination of substantial equivalence is granted on the condition that prior to shipping the first device, you submit a postclearance special report. This report should contain complete information, including acoustic output measurements based on production line devices, requested in Appendix G, (enclosed) of the Center's September 30, 1997 "Information for Manufacturers Seeking Marketing Clearance of Diagnostic Ultrasound Systems and Transducers." If the special report is incomplete or contains unacceptable values (e.g., acoustic output greater than approved levels), then the 510(k) clearance may not apply to the production units which as a result may be considered adulterated or misbranded.

The special report should reference the manufacturer's 510(k) number. It should be clearly and prominently marked "ADD-TO-FILE" and should be submitted in duplicate to:

Food and Drug Administration  
Center for Devices and Radiological Health  
Document Mail Center (HFZ-401)  
9200 Corporate Boulevard  
Rockville, Maryland 20850

This letter will allow you to begin marketing your device as described in your premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus permits your device to proceed to market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (301) 594-4591. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or at (301) 443-6597 or at its Internet address "<http://www.fda.gov/cdrh/dsmamain.html>".

If you have any questions regarding the content of this letter, please contact Rodrigo C. Perez at (301) 594-1212.

Sincerely yours,



*for* Nancy C. Brogdon  
Director, Division of Reproductive,  
Abdominal and Radiological Devices  
Office of Device Evaluation  
Center for Devices and Radiological Health

Enclosure(s)

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **SONOLINE G20™ Diagnostic Ultrasound Systems**

Intended Use: Diagnostic imaging or fluid flow analysis of the human body as follows:

Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		P	P						BM	Note 2,3
Abdominal		P	P						BM	Note 2,3
Intraoperative (Note 6)		P	P						BM	Note 3
Intraoperative Neurological		P	P						BM	Note 3
Pediatric		P	P						BM	Note 2,3
Small Organ (Note 1)		P	P						BM	Note 2,3
Neonatal Cephalic		P	P						BM	Note 3
Adult Cephalic		P	P						BM	Note 2
Cardiac		P	P						BM	Note 2
Transesophageal		P	P						BM	Note 2,3
Transrectal		P	P						BM	Note 2,3
Transvaginal		P	P						BM	Note 2,3
Transurethral										
Intravascular										
Peripheral vessel		P	P						BM	Note 2,3
Laparoscopic		P	P						BM	Note 3
Musculo-skeletal Conventional		P	P						BM	Note 2,3
Musculo-skeletal Superficial		P	P						BM	Note 2,3
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

Note 1 For example: breast, testes, thyroid, penis, prostate, etc.

Note 2 Ensemble tissue harmonic imaging

Note 3 3D imaging

Note 4 B&W SieScape panoramic imaging

Note 5 Power SieScape panoramic imaging

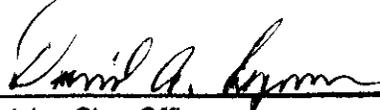
Note 6 For example: abdominal, vascular

Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **C5-2 Convex Array Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**  
 Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

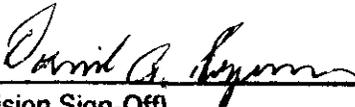
Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		P	P						BM	Note 2,3
Abdominal		P	P						BM	Note 2,3
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric		P	P						BM	Note 2,3
Small Organ										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Trans-esophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral vessel		P	P						BM	Note 2,3
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (Specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **L10-5 Linear Array Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**  
 Intended Use: **Ultrasound imaging or fluid flow analysis of the human body as follows:**

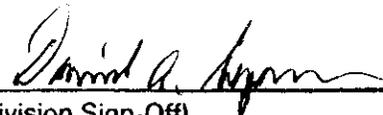
Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal										
Abdominal		P	P						BM	Note 2,3
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric		P	P						BM	Note 2,3
Small Organ		P	P						BM	Note 2,3
Neonatal Cephalic		N	N						BM	Note 2,3
Adult Cephalic										
Cardiac										
Trans-esophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral vessel		P	P						BM	Note 2,3
Laparoscopic										
Musculo-skeletal Conventional		P	P						BM	Note 2,3
Musculo-skeletal Superficial		P	P						BM	Note 2,3
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)



(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **EV9-4 Convex Array Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**  
 Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		P	P						BM	Note 3
Abdominal										
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric										
Small Organ (Note 1)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal		P	P						BM	Note 3
Transvaginal		P	P						BM	Note 3
Transurethral										
Intravascular										
Peripheral vessel										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*David A. Nguyen*  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number           K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **7.5L75S** Linear Array Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems  
 Intended Use: Diagnostic imaging or fluid flow analysis of the human body as follows:

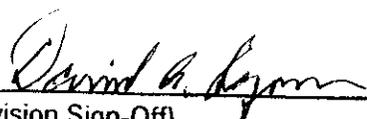
Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal										
Abdominal		P	P							Note 3
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric		P	P							Note 3
Small Organ (Note 1)		P	P							Note 3
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral vessel		P	P							Note 3
Laparoscopic										
Musculo-skeletal Conventional		P	P							Note 3
Musculo-skeletal Superficial		P	P							Note 3
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **EC9-4 Convex Array Endocavity Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**  
 Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

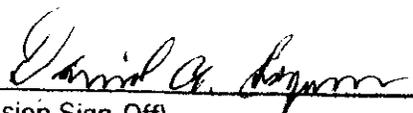
Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		P	P						BM	Note 3
Abdominal										
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric										
Small Organ (Note 1)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal		P	P						BM	Note 3
Transvaginal		P	P						BM	Note 3
Transurethral										
Intravascular										
Peripheral vessel										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **C4-2 Convex Array Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**

Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		P	P						BM	
Abdominal		P	P						BM	Note 2,3
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric		P	P						BM	
Small Organ (Note 1)										
Neonatal Cephalic										
Adult Cephalic		P	P						BM	Note 2,3
Cardiac		P	P						BM	Note 2,3
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral vessel										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*David A. Segon*  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **C8-5 Convex Array Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**  
 Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

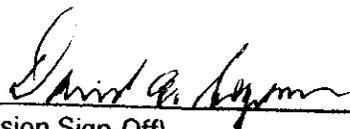
Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal										
Abdominal		P	P						BM	Note 3
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric		P	P						BM	Note 3
Small Organ (Note 1)		P	P						BM	Note 3
Neonatal Cephalic		P	P						BM	Note 3
Adult Cephalic										
Cardiac		P	P						BM	Note 3
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral vessel										
Laparoscopic										
Musculo-skeletal Conventional		P	P						BM	Note 3
Musculo-skeletal Superficial		P	P						BM	Note 3
Other (specify)										

N = new indication; P = previously cleared by FDA with **K040502**; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K04-2833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **BE9-4 Biplane Endocavity Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**

Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

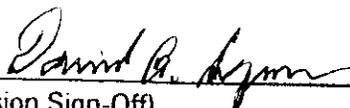
Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		N	N						BM	Note 3
Abdominal										
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric										
Small Organ (Note 1)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal		N	N						BM	Note 3
Transvaginal		N	N						BM	Note 3
Transurethral										
Intravascular										
Peripheral vessel										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **C6F3 Convex Array Mechanically Driven, 3D Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**  
 Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

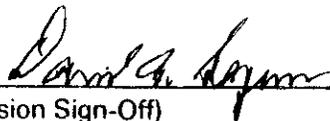
Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		P	P						BM	Note 2,3
Abdominal		P	P						BM	Note 2,3
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric		P	P						BM	Note 2,3
Small Organ (Note 1)										
Neonatal Cephalic		P	P						BM	Note 2,3
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral vessel										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **EV8F5 Mechanical Sector Endovaginal 3D Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**

Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

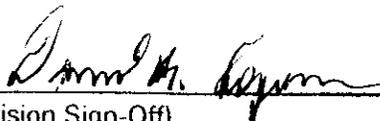
Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		P	P						BM	Note 3
Abdominal										
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric										
Small Organ (Note 1)										
Neonatal Cephalic		P	P						BM	Note 3
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal		P	P						BM	Note 3
Transvaginal		P	P						BM	Note 3
Transurethral										
Intravascular										
Peripheral vessel										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **3.5C40S Convex Array Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**  
 Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		P	P						BM	Note 2,3
Abdominal		P	P						BM	Note 2,3
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric		P	P						BM	Note 2,3
Small Organ (Note 1)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral vessel		P	P						BM	Note 2,3
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*David G. Reymann*  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **5.0C40S Convex Array Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**

Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

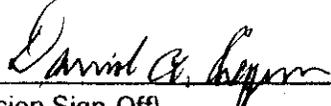
Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		P	P						BM	Note 2,3
Abdominal		P	P						BM	Note 2,3
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric		P	P						BM	Note 2,3
Small Organ (Note 1)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral vessel		P	P						BM	Note 2,3
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N = new indication; P = previously cleared by FDA with K040502; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

  
 (Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number           K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **Endo PII Mechanical Sector Biplane Endo-cavity Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**

Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		N	N						BM	Note 2,3
Abdominal		N	N						BM	Note 2,3
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric		N	N						BM	Note 2,3
Small Organ (Note 1)										
Neonatal Cephalic										
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal										
Transvaginal										
Transurethral										
Intravascular										
Peripheral vessel		N	N						BM	Note 2,3
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N = new indication; P = previously cleared; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)

*David C. Kopyov*

(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number K042833

**Diagnostic Ultrasound Indications for Use Form**

510(k) Number (if known):

Device Name: **Endo VII Mechanical Sector Endovaginal Transducer for use with:  
 SONOLINE G20™ Diagnostic Ultrasound Systems**

Intended Use: **Diagnostic imaging or fluid flow analysis of the human body as follows:**

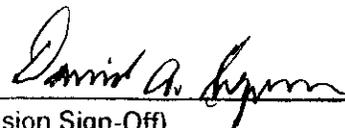
Clinical Application	Mode of Operation									
	A	B	M	PWD	CWD	Color Doppler	Amplitude Doppler	Color Velocity Imaging	Combined (Specify)	Other (Specify)
Ophthalmic										
Fetal		N	N						BM	Note 2,3
Abdominal										
Intraoperative Abdominal										
Intraoperative Neurological										
Pediatric										
Small Organ (Note 1)										
Neonatal Cephalic		N	N						BM	Note 2,3
Adult Cephalic										
Cardiac										
Transesophageal										
Transrectal		N	N						BM	Note 2,3
Transvaginal		N	N						BM	Note 2,3
Transurethral										
Intravascular										
Peripheral vessel										
Laparoscopic										
Musculo-skeletal Conventional										
Musculo-skeletal Superficial										
Other (specify)										

N = new indication; P = previously cleared; E = added under Appendix E

- Note 1 For example: breast, testes, thyroid, penis, prostate, etc.
- Note 2 Ensemble tissue harmonic imaging
- Note 3 3D imaging
- Note 4 B&W SieScape panoramic imaging
- Note 5 Power SieScape panoramic imaging
- Note 6 For example: abdominal, vascular
- Note 7 Contrast agent imaging

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)  
 Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use (Per 21 CFR 801.109)



(Division Sign-Off)  
 Division of Reproductive, Abdominal,  
 and Radiological Devices  
 510(k) Number                     K042833