

Specificați Completată

Model: Logiq P9; Producător: GE Healthcare în colaborare cu GE Ultrasound Korea, GE MEDICAL SYSTEM SCS; Țara: SUA, KOREA și Franța.

Specificarea tehnică deplină solicitată de către autoritatea contractantă	Specificarea tehnică deplină propusă de către autoritatea ofertantă
APLICAȚII CLINICE: General; PORTURI ACTIVE PENTRU TRANSDUCTORI – 3; PORTURI PENTRU TRANSDUCTORI CW – 1; NIVELE DE GRI ≥ 256; GAMA DINAMICĂ ≥ 250dB; PREPROCESARE; CANALE DIGITALE ≥ 2 mil.; ADINCIME DE SCANARE ≥ 30 cm; DIAPAZON FRECVENTĂ ASIGURATĂ DE DISPOZITIV (minim) – 1-18 Mhz; TRANSDUCTORI ACCEPTAȚI DE SISTEM: Lineare matriciale; Convexe matriciale; TEE; Intra-operative; Sectoriale matriciale; MULTIFRECVENTA TRANSDUCTOARE ≥ 8; POSTPROCESARE; IMAGINE MODURI: M-mod; M-mod și 2-D; Armonici Tisulare; Armonici Tisulare diferențiale; M-mode anatomic; M-Mode color; DOPPLER: Tip CW, PW, CFM, TDI; Măsurări automatizate; Calcul automate; Power Doppler; Duplex; Triplex; FUNCȚIONALITĂȚI: Ajustare frecvență; Diapazon dinamic reglabil; Focalizare pe imagine vertical/orizontal – 1-5 zone; Ajustare mape de culori ≥ 9; Selectare automată a sondei la aplicarea presetului; Reglare GAIN; Reglarea semnalului acustic; Măsurători în timp real și în freeze; PAN/ZOOM: Imagine în timp real; Imagine înghețată; STOCARE IMAGINI: Capacitate ≥ 240GB SSD; Memorie CINE ≥ 700MB sau 60; CD/DVD; USB 3.0, 2.0; PACHETE DE ANALIZĂ: General; Muculoschiletal; Vascular; DICOM 3.0; TRANSDUCTOARE: Linear – 5-18 Mhz; Linear – 3-13 Mhz; MONITOR FULL HD ≥ 21";	APLICAȚII CLINICE: General; PORTURI ACTIVE PENTRU TRANSDUCTORI – 4; DA PORTURI PENTRU TRANSDUCTORI CW – 1; DA NIVELE DE GRI ≥ 256; DA GAMA DINAMICĂ ≥ 250dB; DA PREPROCESARE; CANALE DIGITALE - 386.469.; DA ADINCIME DE SCANARE - 33 cm; DA DIAPAZON FRECVENTĂ ASIGURATĂ DE DISPOZITIV – 1-20Mhz; DA TRANSDUCTORI ACCEPTAȚI DE SISTEM: Lineare matriciale; DA Convexe matriciale; DA pînă la 192 elemente TEE; DA Intra-operative; DA Sectoriale matriciale; DA pînă la 96 de elemente MULTIFRECVENTA TRANSDUCTOARE ≥ 8; DA în dependență de sondă POSTPROCESARE; IMAGINE MODURI: M-mod; DA M-mod și 2-D; DA Armonici Tisulare; DA Armonici Tisulare diferențiale; DA M-mode anatomic; DA M-Mode color; DA DOPPLER: Tip CW, PW, CFM, TDI; DA Măsurări automatizate; DA Calcul automate; DA Power Doppler; DA Duplex; DA Triplex; DA FUNCȚIONALITĂȚI: Ajustare frecvență; DA Diapazon dinamic reglabil; DA Focalizare pe imagine vertical/orizontal – 1-5 zone; DA în dependența sondei Ajustare mape de culori ≥ 9; DA Selectare automată a sondei la aplicarea presetului; Reglare GAIN; DA Reglarea semnalului acustic; DA Măsurători în timp real și în freeze; DA PAN/ZOOM: DA Imagine în timp real; DA Imagine înghețată; DA STOCARE IMAGINI: Capacitate 345 GB HDD; DA Memorie CINE 776MB DA; CD/DVD; DA USB 3.0, 2.0; DA PACHETE DE ANALIZĂ: General; DA Muculoschiletal; DA Vascular; DA DICOM 3.0; DA TRANSDUCTOARE: Linear – 7-20 Mhz; DA L10-20-RS Linear – 2-11 Mhz; DA L3-12-RS MONITOR FULL HD – 21,5"; DA

BUOTOANE CONSOLA – Configurabile; TASTATURA DIGITALA; BRAȚ ARTICULAT – flexibil; Posibilitatea efectuării Up Grade; ACCESORII: B/W printer – incorporat; DVD/CD RW – incorporat; GEL USG 1 litru – 10 bucăți; Hirtie USG – 10 bucăți; Cu 4 roți blocabile. Cerințe de certificare: * Certificat de la producător ce atestă calitatea produsului ISO 13485, valabil, copie confirmată prin semnatura și ștampila Participantului. * Certificat CE sau declarație de conformitate CE cu anexele corespunzătoare pentru produsele oferite, valabil, copie confirmată prin semnatura și ștampila Participantului. * Declarație de la Ofertant – confirmată prin semnatura și ștampila, în care să certifice termenul de garanție pentru echipament și accesorii nu mai mic de 36 luni din momentul instalării/dării în exploatare a bunului. *Trăning pentru utilizatori la instalare și la solicitare – obligatoriu. Documente confirmative: *Manual deservice în una din limbile de circulație internațională (rusa/engleza). *Manual de utilizare cu prezentarea traducerii la momentul livrării în limba română, copie, confirmată prin semnătura și ștampila Participantului. *Ghid rapid al utilizatorului (max. 4 pagini A4), în limba de stat - obligatoriu	BUOTOANE CONSOLA – Configurabile; DA TASTATURA DIGITALA; DA BRAȚ ARTICULAT – flexibil; DA Posibilitatea efectuării Up Grade; DA ACCESORII: B/W printer – incorporat; DA DVD/CD RW – incorporat; DA GEL USG 1 litru – 10 bucăți; DA Hirtie USG – 10 bucăți; DA Cu 4 roți blocabile. DA Cerințe de certificare: * Certificat de la producător ce atestă calitatea produsului ISO 13485, valabil, copie confirmată prin semnatura și ștampila Participantului. DA * Certificat CE sau declarație de conformitate CE cu anexele corespunzătoare pentru produsele oferite, valabil, copie confirmată prin semnatura și ștampila Participantului. DA * Declarație de la Ofertant – confirmată prin semnatura și ștampila, în care să certifice termenul de garanție pentru echipament și accesorii nu mai mic de 36 luni din momentul instalării/dării în exploatare a bunului. DA *Trăning pentru utilizatori la instalare și la solicitare – obligatoriu. DA Documente confirmative: *Manual deservice în una din limbile de circulație internațională (rusa/engleza). DA *Manual de utilizare cu prezentarea traducerii la momentul livrării în limba română, copie, confirmată prin semnătura și ștampila Participantului. DA *Ghid rapid al utilizatorului (max. 4 pagini A4), în limba de stat – obligatoriu DA
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ATTESTATION CE / EC CERTIFICATE

Approbation du Système Complet d'assurance Qualité/ Approval of full Quality Assurance System

ANNEXE II excluant le point 4 Directive 93/42/CEE relative aux dispositifs médicaux

ANNEX II excluding section 4 Directive 93/42/EEC concerning medical devices

Pour les dispositifs de classe III, un certificat CE de conception est requis

For class III devices, a EC design certificate is required

Fabricant / Manufacturer

GE ULTRASOUND KOREA, Ltd.

9, Sunhwan-ro 214beon-gil, Jungwon-gu,

SEONGNAM-SI, GYEONGGI-DO, REPUBLIC OF KOREA

Catégorie du(des) dispositif(s) / Device(s) category

Dispositif ou système de diagnostic par ultrasons

Ultrasound diagnostic device or system

Voir document complémentaire GMED / See GMED additional document

n° 36988

GMED atteste qu'à l'examen des résultats figurant dans le rapport référencé P183396, P601203, le système d'assurance qualité - pour la conception, la production et le contrôle final - des dispositifs médicaux énumérés ci-dessus est conforme aux exigences de l'annexe II excluant le point 4 de la Directive 93/42/CEE.

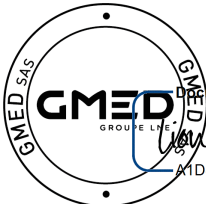
GMED certifies that, on the basis of the results contained in the file referenced P183396, P601203, the quality system - for design, manufacturing, and final inspection - of medical devices listed here above complies with the requirements of the Directive 93/42/EEC, annex II excluding section 4.

La validité du présent certificat est soumise à une vérification périodique ou imprévue

The validity of the certificate is subject to periodic or unexpected verification

Début de validité / Effective date : March 17th, 2021 (included)

Valable jusqu'au / Expiry date : May 26th, 2024 (included)


Signed by:
Lionel DREUX
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Lionel DREUX
Certification Director

Ce document complémentaire GMED n° 36988 rev. 1 atteste de la validité du certificat CE N° 7697 rev. 19 au regard des informations listées ci-dessous.

This GMED additional document n° 36988 rev. 1 attests to the validity of EC certificate N° 7697 rev. 19 with regard to the information listed below.

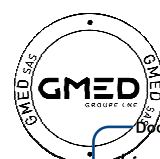
**Fabricant / Manufacturer: GE ULTRASOUND KOREA, Ltd.
9, Sunhwan-ro 214beon-gil, Jungwon-gu,
SEONGNAM-SI, GYEONGGI-DO, REPUBLIC OF KOREA**

Identification des dispositifs / Identification of devices

Désignation du dispositif / Accessoires marqués CE <i>Device designation / CE marked accessories</i>	Réf commerciale du dispositif ou code article <i>Device commercial reference or article code</i>	Classe du DM MD class
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	LOGIQ P7	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	LOGIQ P8	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	LOGIQ P9	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	LOGIQ P10	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	VOLUSON S6	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	VOLUSON S8	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	VOLUSON S8t	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	VOLUSON S10	Ila

GMED 0459

GMED - 36988 rev. 1
Modifie le document n° 36988 rev.0



DocuSigned by:

Lionel DREUX

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Lionel DREUX
Certification Director

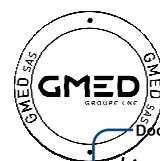
Désignation du dispositif / Accessoires marqués CE <i>Device designation / CE marked accessories</i>	Réf commerciale du dispositif ou code article <i>Device commercial reference or article code</i>	Classe du DM <i>MD class</i>
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	VOLUSON S10 Expert	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	VOLUSON P6	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	VOLUSON P8	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	VOLUSON SWIFT	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	VOLUSON SWIFT+	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	LOGIQ S8	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	LOGIQ S8 T1	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	LOGIQ S7 Expert	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	LOGIQ S7 Pro	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	LOGIQ S7 XDclear2.0	Ila
Dispositif ou système de diagnostic par ultrasons <i>Ultrasound diagnostic device or system</i>	LOGIQ E10s	Ila

Site couvert et Activités / Location and Activities

Site / Location	Activités / Activities
GE ULTRASOUND KOREA, Ltd. 9, Sunhwan-ro 214beon-gil, Jungwon-gu, Seongnam-si, Gyeonggi-do, REPUBLIC OF KOREA équivalent à <i>equivalent to</i> GE ULTRASOUND KOREA, Ltd. 65-1, Sangdaewon-dong, Jungwon-gu, Seongnam-si, Gyeonggi-do - 462-120 REPUBLIC OF KOREA	Conception, fabrication et contrôle final <i>Design, manufacture and final control</i>

GMED 0459

GMED - 36988 rev. 1
Modifie le document n° 36988 rev.0



DocuSigned by:

Lionel DREUX

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Lionel DREUX
Certification Director



EC Declaration of Conformity

Following the provisions of the medical devices directive 93/42/EEC, Annex II and of the directive 2011/65/EU, directive 2012/19/EU, directive 2014/53/EU

Manufacturer:

**GE Ultrasound Korea, Ltd.
9, Sunhwan-ro 214beon-gil,
Jungwon-gu, Seongnam-si,
Gyeonggi-do Republic of Korea**

EU Authorized Representative:

**GE MEDICAL SYSTEMS SCS
283 RUE DE LA MINIERE
78530 BUC
FRANCE**

Equivalent to

**65-1, Sangdaewon-dong,
Jungwon-gu, Seongnam-si
Gyeonggi-do 462-120 Republic of Korea**

*We hereby declare under our sole responsibility that the class **Ila** product:*

LOGIQ P7, LOGIQ P9, General Purpose Ultrasound Imaging System including accessories and components (ref: See Addendum)

GMDN Code: **40761**

UMDNS Code: **15976**

Classification rule (93/42/EC Annex IX): **Rule 10**

To which this declaration relates, is in conformity with the requirements of:

The medical devices directive 93/42/EEC (MDD)

The directive 2011/65/EU on the restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS)

The directive 2012/19/EU on the waste electrical and electronic equipment (WEEE)

The directive 2014/53/EU on the radio equipment (RED)

The Commission Regulation (EU) No 207/2012 of 9 March 2012 on electronic instructions for use of medical devices

This conformity is based on the following elements:

- Information included in the technical documentation ref.: **DOC1587707** /DHF ref.: **DOC1412680**, of the product to which this declaration relates.
- EC certificate: approval of full quality assurance system (Annex II of the medical devices directive 93/42/EEC) delivered by **GMED** (Notified Body N° **0459**) on Certificate Number **N° 7697**.



▪ List of harmonized standards applied for CE marking

- EN 60601-1:2006/A12:2014 (Edition 3.1)
- EN 60601-1-2:2015
- EN 60601-1-6:2010/A1:2015
- EN 60601-2-37: 2008/A1:2015
- EN 62304:2006/AC: 2008
- EN 62366:2008 + A1:2015
- EN 1041:2008
- EN ISO 15223-1: 2016

This EC declaration of conformity supersedes the previous declaration dated **26-Mar-2019**.

A handwritten signature in blue ink, appearing to read 'Soyoung Park', written over a horizontal line.

Park, Soyoung
Regulatory Affairs Specialist

Date/Datum: 17-May-2019

GE Healthcare. GE Ultrasound Korea, Ltd.
9, Sunhwan-ro 214beon-gil, Jungwon-gu, Seongnam-si,
Gyeonggi-do Republic of Korea



ADDENDUM TO THE EC DECLARATION OF CONFORMITY dated 17-May-2019

- **GTIN#:**
 - ✓ LP7 R3 – 00840682141314
 - ✓ LP9 R3 – 00840682141246

Product Description	HCAT #
Base Systems	
LOGIQ P7 R3	H42872LA
LOGIQ P9 R3	H42872LB
Probes	
3Sc-RS Probe	H45041DL
6S-RS PROBE	H45021RP
12S-RS Probe	H44901AB
ML6-15-RS Probe	H40462LM
L3-12-RS Probe	H44901AP
L4-12t-RS Probe	H48062AB
L12n-RS probe	H48062AH
12L-RS Probe	H40402LY
L6-12-RS Probe	H48062AC
9L-RS Probe	H40442LL
L10-22-RS Probe	H48312AH
L8-18i-RS Probe	H40462LF
C1-5-RS Probe	H40462LA
4C-RS Probe	H4000SR
8C-RS Probe	H40402LS
E8C-RS Probe	H40402LN
E8Cs-RS Probe	H48062AF
IC9-RS Probe	H48691PJ
BE9CS-RS Probe	H40482LN
RAB2-6-RS Probe	H48681WR
RIC5-9A-RS Probe	H48701EJ
6Tc-RS Probe	H45551ZE
Doppler P8D Probe	H46312LZ
P6D	H4830JG
P2D	H4830JE
L3-9i-RS Probe	H46442LK
Biopsy Options	
3SP MULTI-ANGLE BIOPSY	H46222LC
9L BIO GUIDE STARTER KIT	H4906BK
12L-RS Biopsy Starter Kit	H40432LC
ML6-15 Biopsy Starter Kit	H40432LJ
12L TRANSVERSE BRACKET	H48392LL
INFINITE 12L BIOPSY KIT	H48392LT



L3-12-D Biopsy starter kit	H48302AA
C1-5 Biopsy Starter Kit	H40432LE
4C BIOPSY BRACKET	E8385NA
E721 STARTER KIT	E8385MJ
E8C E721 E8C-RS IC5-9H MTZ Biopsy Kit	E8333JB
E8C REUSABLE BIOPSY KIT	H40412LN
BE9CS Biopsy Kit 742-339	H42742LH
BE9CS Biopsy Kit 742-401	H42742LJ
Reusable Biopsy Needle Guide for GE BE9C Ultrasound Probe	E8387MA
Sterile Disposable Biopsy Needle Guide Kit for GE BE9C Probe	E8387M
IC9 reusable Biopsy	H48701MN
IC9 Biopsy Starter Kit	H48691YW
RAB6-D BIOPSY STARTER KIT	H48681ML
PEC63 BIOPSY KIT FOR RIC5-9	H46721R
RIC STERILE NEEDLE GUIDE	H48681GF
TEE accessory	
TEE Cleaning and Storage System	H45551NK
TEE Storage Rack	H45551NM
TEE Scan Head Protection Cover	H45521CK
TEE Clip-on Bite Guard Adult	H45511EE
TEE Clip-on Bite Guard Adult OR	H45521CB
Conventional Bite Guard Adult	H45521JH
Bite Hole Indicator	H45531HS
Software options	
LP7 and LP9 Advanced 3D	H42782LK
LP7 and LP9 Auto IMT	H42782LL
LP7-P9 R3 HD B-Flow	H42892LR
LP7-P9 R3 CEUS	H42892LS
LP7-P9 R3 HRes CEUS	H42892LT
LP7 and LP9 DICOM	H42782LR
LP7 and LP9 Elastography	H42782LS
LP7 and LP9 Elastography Quantification	H42782LT
LP7 and LP9 Flow Quantification	H42782LW
LP7 and LP9 LOGIQView	H42782LY
LP7 and LP9 Report Writer	H42782LZ
LP7 and LP9 Scan Assistant	H42792LA
LP7 and LP9 Stress Echo	H42792LB
LP7 and LP9 Tissue Velocity Imaging TVI	H42792LC
LP7 and LP9 B Steer+	H42792LD
LP7 and LP9 4D TUI Software	H42792LF
LP7 and LP9 VOCAL Software	H42792LG
LP7 and LP9 VCI Static Software	H42792LH
LP7-P9 STIC	H42822LZ
LP7-P9 Omniview	H42832LA



LP7-P9 R3 HDLive	H42892LW
LP7 and LP9 Auto EF	H42792LJ
LP7 and LP9 Meas Assist Breast	H42792LK
LP7 and LP9 Meas Assist OB	H42792LL
LP7 and LP9 Breast Prod	H42792LM
LP7 and LP9 Compare Assistant	H42792LN
LP7 and LP9 Thyroid Prod	H42792LP
LP7 and LP9 SWDVR	H42792LR
LP7 MSK Korea	H42762LF
LP7-P9 Cardiac Strain	H42822LY
LP7-P9 R2.5 Pinpoint GT option	H40292LC
LP7-P9 R3 Shear Wave Elastography	H42892LY
LOGIQ P Apps	H42892LZ
LOGIQ P Apps without Dongle	H42922LM
LOGIQ P7 R3 Advanced Probes Convex	H42922LN
LOGIQ P7 R3 Advanced Probes Linear	H42922LP
LOGIQ P7 R3 Advanced Probes Sector	H42922LR
LOGIQ P7 R3 Advanced Probes Specialty	H42922LS
Hardware options	
LP7-P9 R3 Card Reader Mounting Kit	H42792LZ
Art. Monitor Arm white	H42902LB
LP7-P9 R3 Rear handle	H42902LC
LP7 and LP9 OPIO tray	H42802LG
LP7 and LP9 Paper tray	H42802LE
LP7 and LP9 Side Tray	H42802LC
LP7-P9 R3 Cable Hook rear	H42902LD
LP7-P9 R3 Gel Warmer	H42902LE
LP7-P9 R3 4 port kit	H42912LF
LP7 and LP9 4D Kit	H42802LD
LP7 P9 CW HW Kit	H46432LN
LP7 P9 Pencil CW HW Kit	H42802LB
USB FOOTSWITCH 3 BUTTON	H46732LF
LP7 P9 W. LESS LAN KIT-J	H42812LD
LP7 P9 W. LESS LAN KIT	H42802LL
LP7 P9 UVC	H42832LJ
LP7 P9 UVC for Japan	H42832LK
ISOLATION TRANSFORMER	H48671WN
Pwr supply noise filter	H46162LH
Pinpoint GT Practice kit	H48672AB
Barcode reader	H48872LG
LP7-P9 R3 ODD Option	H42912LE
ECG options	
LP7 P9 ECG module only	H42792LS
LS8 ECG CABLE - AHA	H46102LW



LS8 ECG CABLE - IEC	H45302LZ
Veterinary Use Only	
Vet kit	H46832LC
Probe Vet Label	H48992LR
Vet probe caution label	H48492AW
Peripherals	
Printers	
UP-D898 BW Printer Kit	H46992LS
LP7 LP9 BW INSTALL KIT	H46432LP
UP-D25MD PRINTER	H44642LW
Cabinet	
LP7-P9 R3 HIGH CABINET	H42902LG
LP7-P9 R3 DRAWER	H42902LH
LP7-P9 R3 Low Cabinet	H42902LJ
Accessories	
LP7-P9 R3 Multi P. holder	H42902LK
LOGIQ S7 R3 Small Probe Holder	H46302LB
PROBE CABLE HANGER	H44412LA
Batteries	
LP7-P9 R2 Battery option	H42832LG
LP7-P9 R3 ext battery	H42902LM
Power Cords	
POWER CORD FIX BRKT 220V	H42812LJ
POWER CORD FIX BRKT 110V	H42812LK
Destination Sets	
DESTINATION SET UK	H46712LM
DESTINATION SET S AFRICA	H46712LN
DESTINATION SET ARGENTINA	H46712LP
DESTINATION SET ISRAEL	H46712LR
DESTINATION SET SWISS	H46712LS
DESTINATION SET DENMARK	H46712LT
DESTINATION SET US	H46712LW
DESTINATION KIT AUS_NZ	H46712LZ
DESTINATION SET CHINA	H46722LA
DESTINATION SET INDIA	H46722LB
DESTINATION SET ITALY	H46722LD
DESTINATION SET BRAZIL	H46752LW
DESTINATION SET Taiwan	H44512LY
DESTINATION SET JAPAN	H46712LY
Keyboards and Key Cap Language Kits	
AN Keyb. Greek black	H42902LR
AN Keyb. Norwegian black	H42902LS
AN Keyb. Russian black	H42902LT
AN Keyb. French black	H42902LW



AN Keyb. Swedish black	H42902LY
AN Keyb. German black	H42902LZ
AN Keyb. English black	H42912LA
Upgrade kit	
LOGIQ P9 R2.5 to R3 SW conversion	H42922LK
LOGIQ P7 R2.5 to R3 SW conversion	H42922LL

Notes:

[1] Catalog number identifies the device(s) in the manufacturer's catalog and is usually included on commercial documents like sales contract, order processing documents and shipping documents.

[2] Probes and accessories may carry the CE-mark and when applicable, the Notified Body number corresponding to the EC Declaration under which the products are CE-marked by their manufacturer. GE Ultrasound Korea Ltd. has verified the mutual compatibility of the devices in combination with LOGIQ P9 and LOGIQ P7 and included relevant information to users with the LOGIQ P9 and LOGIQ P7 instructions for use.

End of Document



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
BS-MDR-099



Product Service

EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III
(Class IIa and Class IIb Devices)

No. G10 075707 0078 Rev. 00

Manufacturer:

GE Healthcare Austria GmbH & Co OG

Tiefenbach 15
4871 Zipf
AUSTRIA

The Certification Body of TÜV SÜD Product Service GmbH certifies that the manufacturer has established, documented and implemented a quality management system as described in Article 10 (9) of the Regulation (EU) 2017/745 on medical devices. Details on device categories covered by the quality management system are described on the following page(s).

The Report referenced below summarises the result of the assessment and includes reference to relevant CS, harmonized standards and test reports. The conformity assessment has been carried out according to Annex IX Chapter I and III of this regulation with a positive result.

The quality management system assessment was accompanied by the assessment of technical documentation for devices selected on a representative basis.

The certified quality management system is subject to periodical surveillance by TÜV SÜD Product Service GmbH. The surveillance assessment shall also include an assessment of the technical documentation for the device or devices concerned on the basis of further representative samples.

Report No.: 713175299

Preceding certificate No.: this certificate is issued for the first time

Valid from: 2020-05-14

Valid until: 2025-05-13

Date of initial issuance / Rev.00: 2020-05-13

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2020-05-14



Benannt durch/Designated by
Zentralstelle der Länder
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bei Arzneimitteln und
Medizinprodukten
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BS-MDR-099



Product Service

EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III
(Class IIa and Class IIb Devices)

No. G10 075707 0078 Rev. 00

Device Group
Echographic Instruments

Risk Classification
IIa

**The validity of this certificate
depends on conditions and/or
is limited to the following:** None

**Revision History including
Changes:** 00 / 2020-05-13 / 713175299



REGISTRUL DE STAT AL DISPOZITIVELOR MEDICALE

Tip	Denumire
I.3. Certificatul CE	Certificat CE
I.2. Declarația de conformitate CE	Declarație de conformitate CE

Nr	Denumire	Den.comerc.	Model	Nr. catalog	Tara	Producatorul	Reprezentant	Ordin	Data	Cod vamal
			logiq p9							
DM000203113	ULTRASONOGRA		LOGIQ P9 R3	H42872LB	Coreea Sud	GE ULTRASOUND KOREA, LTD.	INTERMED S.R.L.	A07.PS-01.Rg04-63	13-03-2019	
DM000131475	ULTRASONOGRA		LOGIQ P9 R1 LATAM, R1 Config.	H42812LN	Austria	GE HEALTHCARE AUSTRIA GMBH & CO OG	INTERMED S.R.L.	A07.PS-01.Rg04-167	25-06-2018	
DM000131376	ULTRASONOGRA		LOGIQ P9, R1 Config.	H46432LM	Austria	GE HEALTHCARE AUSTRIA GMBH & CO OG	INTERMED S.R.L.	A07.PS-01.Rg04-167	25-06-2018	
DM000131473	ULTRASONOGRA		LOGIQ P9 R1 ASEAN, R1 Config.	H46432LW	Austria	GE HEALTHCARE AUSTRIA GMBH & CO OG	INTERMED S.R.L.	A07.PS-01.Rg04-167	25-06-2018	
DM000131482	ULTRASONOGRA		LOGIQ P9 R2, R2 Config.	H46442LM	Austria	GE HEALTHCARE AUSTRIA GMBH & CO OG	INTERMED S.R.L.	A07.PS-01.Rg04-167	25-06-2018	
DM000131602	ULTRASONOGRA		LOGIQ P9 R2.5, R2.5 Config.	H40282LT	Austria	GE HEALTHCARE AUSTRIA GMBH & CO OG	INTERMED S.R.L.	A07.PS-01.Rg04-167	25-06-2018	

 Содержит([Model], 'logiq.p9')

ОЧИСТИТЬ

Certificate

The Certification Body of
TÜV Rheinland LGA Products GmbH

hereby certifies that the organization

GE ULTRASOUND KOREA, Ltd.
9, Sunhwan-ro 214beon-gil, Jungwon-gu
SEONGNAM-SI, GYEONGGI-DO
Republic of Korea

has established and applies a quality management system for medical devices
for the following scope:

(see attachment for scope and additional site included)

Proof has been furnished that the requirements specified in

EN ISO 13485:2016

are fulfilled. The quality management system is subject to yearly surveillance.

Effective Date: 2020-03-17
Certificate Registration No.: SX 60146260 0001
An audit was performed. Report No.: 32090188 001
This Certificate is valid until: 2021-11-04

Certification Body



Deutsche
Akkreditierungsstelle
D-ZM-14169-01-02

Date 2020-03-17



Balazs Bozsik

Balazs Bozsik

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
Tel.: +49 221 806-1371 Fax: +49 221 806-3935 e-mail: cert-validity@de.tuv.com <http://www.tuv.com/safety>

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

**Attachment to
Certificate**

Registration No.: SX 60146260 0001
Report No.: 32090188 001

Organization: GE ULTRASOUND KOREA, Ltd.
9, Sunhwan-ro 214beon-gil, Jungwon-gu
SEONGNAM-SI, GYEONGGI-DO
Republic of Korea

Scope: Design and Development, Manufacture and Final Test of
Ultrasound Diagnostic Devices and Systems

Site Included:
GE Ultrasound Korea, Ltd.
65-1, Sangdaewon-dong, Jungwon-gu
Seongnami-si, Gyeonggi-do
462-120 Republic of Korea

Design and Development, Manufacture and Final Test of
Ultrasound Diagnostic Devices and Systems

Certification Body



Balk Balazs

Date: 2020-03-17

Balazs Bozsik



LOGIQ™ P9

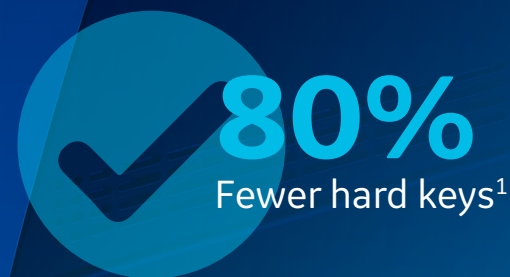
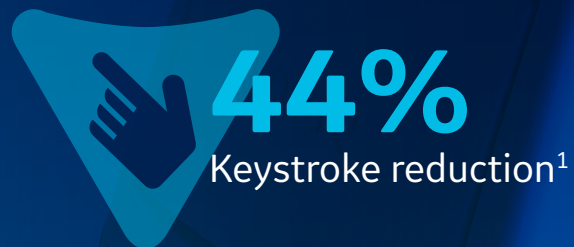
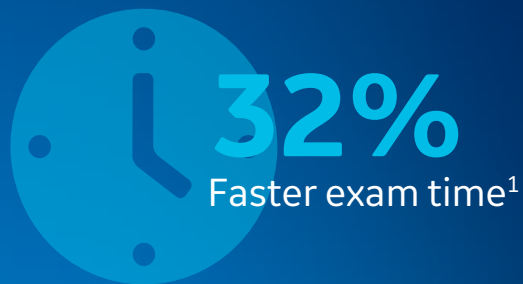
MAKE IT EASY. MAKE IT YOUR OWN.





YOU ASKED FOR SIMPLICITY. **We delivered.**

The LOGIQ™ P9 ultrasound system is ideal for the clinical and workflow demands of general imaging. From triage to comprehensive exams, the budget-friendly LOGIQ P9 delivers consistent image quality, comprehensive application coverage, and ease of use that enable timely, confident decisions.



PERSONALIZED for customized workflow

Easy to learn and use, the LOGIQ P9 supports high efficiency for busy schedules.

Touch Control – Manipulate images on the touch panel with your fingertips, even when gloved. Zoom, magnify, freeze—all with a touch, similar to using a tablet or smartphone. Excellent cleanability as well.

My Page – Personalized digital user interface enables customization of workflow preferences and use case presets. Simply log in, select the appropriate exam type, and begin scanning.

Photo Assistant App – Integrates anatomical photos, taken from an Android™ device, into the ultrasound study to assist clinicians in confirming findings, documenting clinical symptoms, and reporting.



PATIENT-CENTRIC for excellent care

The LOGIQ P9 supports your diagnostic confidence across a wide range of patient exams.

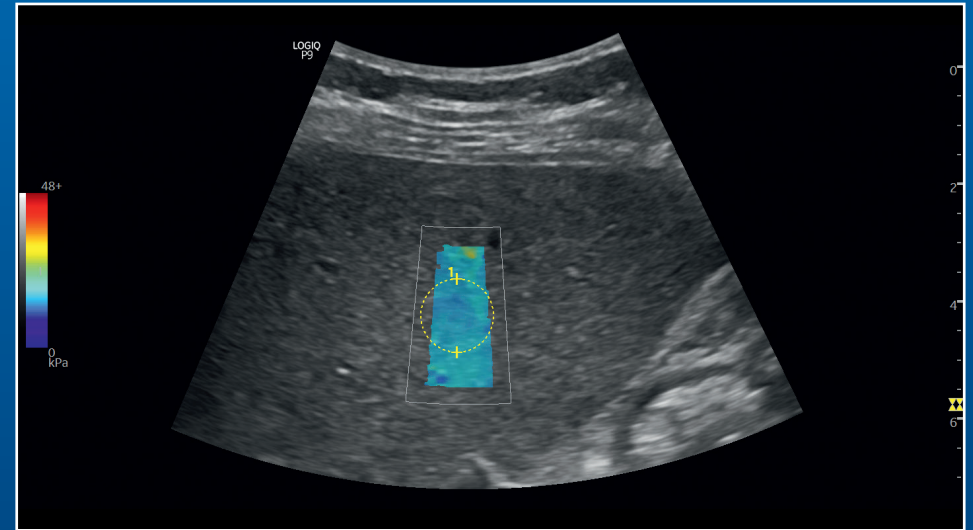
Excellent image quality with minimal tweaking required, and superb B-Mode spatial and contrast resolution.

Wide selection of high quality probes for excellent exam coverage including abdominal, cardiac, OB/GYN, musculoskeletal, vascular, small parts, pediatrics, urology and intra-operative procedures.

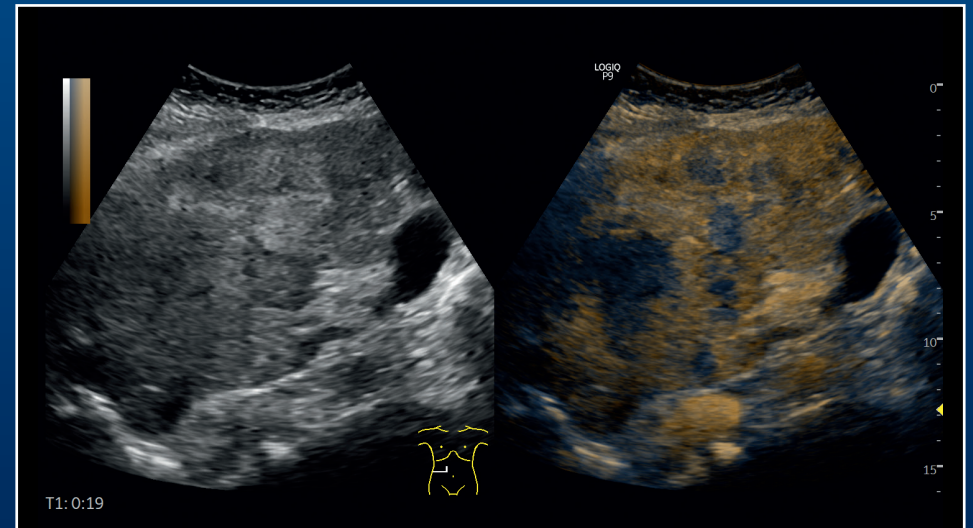
Advanced imaging and visualization tools, including:

- B-Flow™
- HD Color
- 3D/4D
- HDlive™
- STIC/Omniview
- Strain and 2D Shear Wave Elastography
- Stress Echo
- TVI/TVD
- Cardiac Strain
- CEUS
- Enhanced B-Steer+
- Simultaneous display for urology

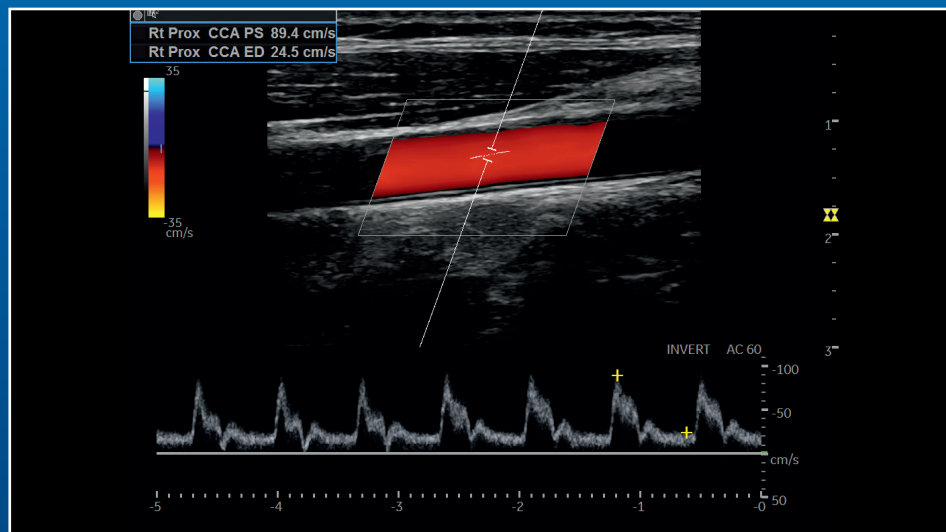
Easily share and archive images with Tricefy™ cloud-based storage, especially useful for OB imaging.



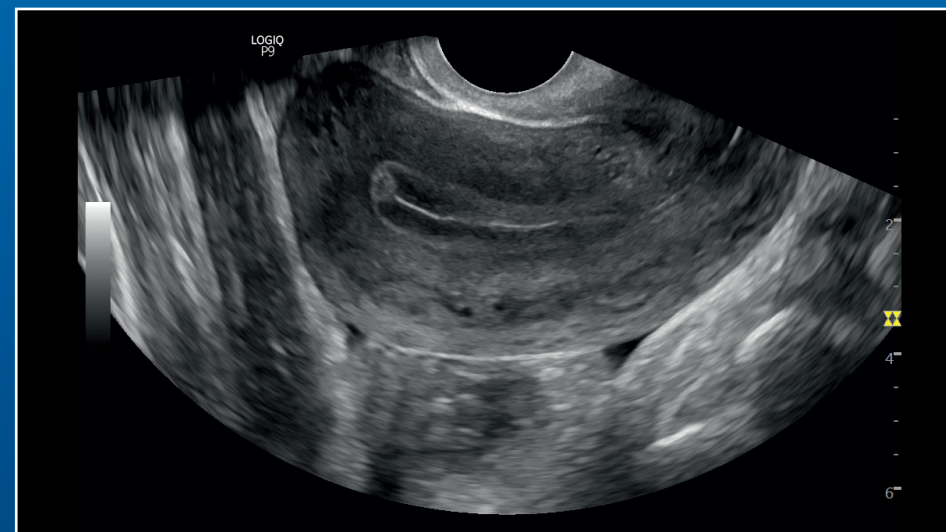
Liver Shear Wave Elastography, C1-5-RS



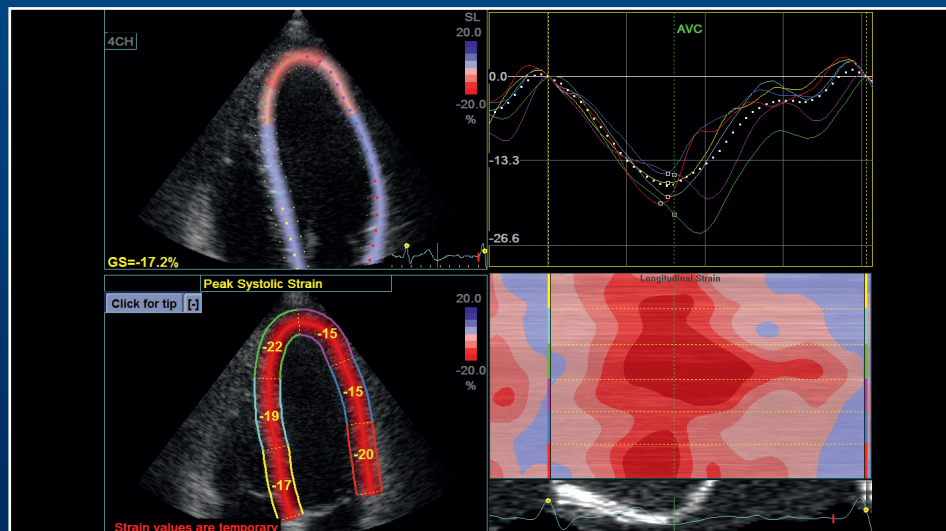
CEUS, C1-5-RS



Carotid Artery CF and PW, L4-12t-RS



Uterus B-Mode, IC9-RS



Cardiac Strain, 3Sc-RS



HDlive, RAB2-6-RS

PRACTICAL for everyday and investment value

The LOGIQ P9 system easily adjusts to your working style, preferences and variable needs throughout your workday.



PRE-PROGRAMMED BUTTON PROBE



REMOTE CONTROL SCANNING

AUTOMATED EFFICIENCY

- Remote Control App enables operation from an Android™ device with the LOGIQ Apps
- Automated tools include Auto TGC, Auto IMT, AutoEF, Measure Assistant, Compare Assistant, and Scan Assistant
- Pre-programmed L4-12t-RS button probe functions without accessing the touch panel to help preserve the sterile field

DATA MANAGEMENT

- Easy data migration supported by DICOM®
- Volume Navigation Import to merge real-time ultrasound with CT and MR datasets

ADVANCED ERGONOMICS

- + External battery enables up to one hour of offline scanning
- + Large 21.5-inch monitor and accessible 10.4-inch touchscreen
- + Simple operating panel design with fewer physical keys¹
- + Fully articulating arm², with up/down/swivel
- + Compact, lightweight design including wireless LAN and Power Assistant battery operation
- + Digital TGC and digital keyboard³

STRONG SECURITY AND SUPPORT

- SonoDefense built on the Windows® 10 IoT operating system provides multi-layer security to protect system integrity and patient data privacy
- My Trainer on-board training modules help accelerate operational confidence
- LOGIQ Club website offers educational resources to help optimize system utilization
- Purchasable service agreement options² including remote service capabilities



Now even more ADVANCED



Touch Control – Easily adjust imaging parameters on touch panel, even wearing gloves



Photo Assistant App – Combine anatomical photos and images in same report



HD Color – Sensitivity for visualizing small vessels and slow flow



2D Shear Wave Elastography – Quantitative estimate of tissue elasticity displayed in color-coded elastograms



Remote Control App – Operate the system from an Android phone or tablet



SonoDefense – Powerful data security features to help guard against costly breaches



Personalized

Customizable workflow features deliver high efficiency your way

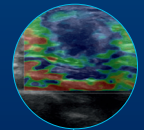


Practical

Advanced automation and ergonomics enable fast exams and reliable results

Patient-centric

Quality images and advanced tools support diagnostic confidence across a wide range of patient exams



The LOGIQ P9 has been designed for compatibility with most commercially available ultrasound contrast agents. Availability of these agents is subject to government regulation and approval. Contrast imaging should be performed within the approved indications for use of the contrast agent used in the exam.

Imagination at work

Product may not be available in all countries and regions. Full product technical specification is available upon request. Contact a GE Healthcare Representative for more information. Please visit www.gehealthcare.com/promotional-locations.

Data subject to change.

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September 2018 JB60493XX 300-19-U001E

1. Internal GE engineering study using standardized protocols for an abdominal exam compared with prior version GE LOGIQ P6 ultrasound system.
2. Availability varies; please contact your GE Healthcare Representative for more information.
3. Optional physical keyboard.



LOGIQ P9

Make it easy. Make it your own.

Product description

The LOGIQ™ P9 is a workhorse for the demanding physician. Its flagship imaging engine is the foundation for finding the root of the patient's problem, even in difficult patients. Buttons on the transducer turn three-handed procedures into two-handed procedures, giving the physician more control. It all adds up to a system that's walk-up easy-to-use on day one and for the most challenging procedures.

The Pinpoint™ GT Needle Guidance Technology is used under license from C.R. Bard, Inc.



General Specification

Dimensions and Weight

Height	Articulating monitor arm 1320 mm ~ 1570 mm (52.0 in ~ 61.8 in)
Width	Keyboard: 430 mm (16.9 in) Foot cover: 495 mm (19.5 in) Monitor: 525 mm (20.7 in)
Depth	Foot cover: 685 mm (27.0 in) Rear handle: 740 mm (29.1 in)
Weight (max. load)	83 kg/183 lbs
Weight (min. load)	68 kg/150 lbs

Electrical Power

Voltage	100 – 240 Vac
Frequency	50/60 Hz
Power consumption maximum of 500 VA with peripherals	

Console Design

4 active probe ports
Integrated solid state drive
Integrated DVD multi-drive (option)
On board storage for BW printer
Integrated speakers
Probe holders
Front handle
Gel warmer (option)
Rear handle (option)
Probe light

User Interface

Operator Keyboard

Ergonomic full size keyboard
Swivel-adjustable, height-adjustable
Digital TGC and digital A/N keyboard
Physical A/N keyboard (option)
10.4" LCD touch screen

Monitor

21.5" widescreen LCD with high resolution

System Overview

Applications

Abdominal
Obstetrical
Gynecological
Breast
Small parts
Musculoskeletal
Vascular
Urological
Pediatric & neonatal
Intraoperative
Cardiac
Transcranial
Endocavitary (transvaginal, transrectal)
Transesophageal

Scanning Methods

Electronic sector
Electronic convex
Electronic micro convex
Electronic linear
Real-time 4D volume sweep

Transducer Types

Sector phased array
Convex array
Microconvex array
Linear array
Matrix array
Single CW (pencil) probes
Volume probes (4D)

Operating Modes

B-Mode
Coded harmonic imaging
M-Mode
Color Flow Mode (CFM)
Power Doppler Imaging (PDI)

System Overview *(cont.)*

Operating Modes *(cont.)*

PW Doppler with high PRF

M-Color Flow Mode

Anatomical M-Mode

Curved anatomical M-Mode

B-Flow™/B-Flow color (option)

Extended Field of View (LOGIQView Option)

Coded Contrast Imaging² (option)

CW Doppler Mode (option)

TVI Mode (option)

Strain Elastography (option)

3D/4D Volume Modes (option)

Shear Wave Elastography (option)

HDlive™ (option)

System Standard Features

Advanced user interface with high resolution 10.4" wide LCD touch screen

Automatic optimization

CrossXBeam™ compounding

Speckle Reduction Imaging (SRI-HD)

Fine angle steering

Coded harmonic imaging

Virtual convex

Advanced 3D (option)

Patient information database

Image archive on integrated CD/DVD (option) and SSD

Raw data analysis

Real-time automatic doppler calculations

OB calculations

Fetal trending

Email to MMS

Mytrainer+

Privacy and security

Qpath

Tricefy™

Multigestational Touch control

System Standard Features *(cont.)*

InSite™ capability

IOTA (International Ovarian Tumor Analysis) LR2 worksheet

Vnav Import

System Options

Auto IMT

Advanced 3D

Cable hook rear

Card reader mounting kit

Strain Elastography

Elastography Quantification³

DICOM® 3.0 connectivity

LOGIQView

B-Flow/B-Flow Color

CF/PDI quantification (FlowQA)

Breast productivity package

Thyroid productivity package

Measure assist OB

AutoEF

B Steer+

Stress echo

Tissue Velocity Imaging (TVI) with Q-Analysis

Scan assistant

Compare assistant

Report writer

Cardiac strain

STIC

OmniView

Guidance Technology (Pinpoint™ GT Needle Guidance Technology)

Shear Wave Elastography

LOGIQ P apps

HDlive

Coded Contrast (CEUS)

HRES CEUS

System Overview *(cont.)*

Peripheral Options

Integrated options for	• Digital BW thermal printer • HDMI output available for compatible devices • S-Video output available for compatible devices • Wireless LAN card for wireless data transfer • External USB printer connection • Power Assistant (battery or extended battery option) for offline scanning
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Digital color thermal printer

Foot switch with programmable functionality

Universal video converter

Barcode reader (for reading needle information)

LOGIQ P apps (Bluetooth)

Display Modes

Live and stored display format: full size and split screen – both with “thumbnails” for still and Cine

Review image format: 4x4 and “thumbnails” for still and Cine

Simultaneous capability

B or CrossXBeam/PW

B or CrossXBeam/CFM or PDI

B/M

B/CrossXBeam

Real-time Triplex Mode (B or CrossXBeam + CFM or PDI/PW or CW (option))

Selectable alternating modes

B or CrossXBeam/PW

B or CrossXBeam + CFM (PDI)/PW(CW (option))

B/CW (option)

Multi-image (split/quad screen)

Live and/or frozen

B or CrossXBeam + B or CrossXBeam/CFM or PDI

Independent Cine playback

Timeline display

Independent dual B or CrossXBeam/PW display

CW

Display formats	• Top/bottom selectable format • Side/side selectable format
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Display Modes *(cont.)*

Virtual convex

Timeline only

Display Annotation

Patient name: first, last and middle

Patient ID

Alternate patient ID

Age, sex and birth date

Hospital name

Date format: 3 types selectable	• MM/DD/YY • DD/MM/YY • YY/MM/DD
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Time format: 2 types selectable	• 24 hours • 12 hours
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Gestational age from	• LMP • EDD	• GA • BBT
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Displayed acoustic output	• TIS: Thermal Index Soft Tissue • TIC: Thermal Index Cranial (Bone) • TIB: Thermal Index Bone • MI: Mechanical Index
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% of maximum power output

Probe name

Map names

Probe orientation

Depth scale marker

Lateral scale marker

Focal zone markers

Image depth

Zoom depth

B-Mode

Gain

Dynamic range

Imaging frequency

Frame averaging

Acoustic frame rate

Gray map

SRI-HD

M-Mode

System Overview *(cont.)*

Display Annotation *(cont.)*

Gain

Dynamic range

Time scale

Doppler mode

Gain

Angle

Sample volume depth and width

Wall filter

Velocity and/or frequency scale

Spectrum inversion

Time scale

PRF

Doppler frequency

Color Flow Mode

Line density

Frame averaging

Packet size

Color scale: 3 types	• Power • Directional PDI • Symmetrical velocity imaging
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Color velocity range and baseline

Color threshold marker

Color gain

PDI

Inversion

Doppler frequency

TGC curve

Cine gage, image number/frame number

Body pattern: multiple human and animal types

Application name

Measurement results

Operator message

Biopsy guide line and zone

Heart rate

General System Parameters

System Setup

Pre-programmable categories

User programmable preset capability

Factory default preset data

Languages: English, French, German, Spanish, Italian, Portuguese, Russian, Greek, Swedish, Danish, Dutch, Finnish, Norwegian, Japanese (message only), Chinese (message only)

OB report formats including Tokyo Univ., Osaka Univ., USA, Europe, and ASUM

User defined annotations

Body patterns

Customized comment home position

Reset

Complete User Manual Available On-Board Through Help (F1)

User manual and service manual are included on USB with each system. A printed manual is available upon request.

CINE Memory/Image Memory

776 MB of Cine memory

Selectable cine sequence for Cine review

Prospective Cine mark

Measurements/calculations and annotations on Cine playback

Scrolling timeline memory

Dual image Cine display

Quad image Cine display

Cine gauge and Cine image number display

Cine review loop

Cine review speed

Image Storage

On-board database of patient information from past exams

Storage formats	• DICOM – compressed/uncompressed, single/multiframe, with/without raw data • Export JPEG, JPEG2000, WMV, MPEG 4 and AVI formats
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Storage devices	• USB memory Stick: 64 MB to 4 GB (for exporting individual images/clips) • CD-R storage: 700 MB • DVD storage: -R (4.7 GB) • Solid state drive image storage: ~345 GB
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General System Parameters *(cont.)*

Image Storage *(cont.)*

Compare old images with current exam

Reload of archived data sets

Connectivity & DICOM

Ethernet network connection

DICOM 3.0 (option)

Wireless LAN (option)

Verify

Print

Store

Modality worklist

Storage commitment

Modality Performed Procedure Step (MPPS)

Media exchange

Off network/mobile storage queue

Query/retrieve

Public SR template

- Structured reporting – compatible with vascular and OB standard
- Direct export DICOM SR and XML

Remote capability InSite™ ExC

DICOM directory import

LOGIQ P apps

Physiological Input Panel (Option)

Physiological input

ECG, 2 lead

Dual R-Trigger

Pre-settable ECG R delay time

Pre-settable ECG position

Adjustable ECG gain control

Automatic heart rate display

Report Writer (Option)

On-board reporting package automates report writing

Formats various exam results into a report suitable for printing or reviewing on a standard PC

Report Writer (Option) *(cont.)*

Exam result reports can include patient info, exam info, measurements, calculations, images, comments and physician diagnosis

Standard templates provided

Customizable templates

Thyroid reporting template

Scanning Parameters

Displayed imaging depth: 0 – 33 cm

Minimum depth of field: 0 – 2 cm (zoom) (probe dependent)

Maximum depth of field: 0 – 33 cm (probe dependent)

Continuous dynamic receive focus/continuous dynamic

Receive aperture

Adjustable dynamic range

Adjustable Field of View (FOV)

Image reverse: right/left

Image rotation of 0°, 90°, 180°, 270°

Digital B-Mode

Adjustable	• Acoustic power • Dynamic range • Gray scale map • Line density • B colorization • Reject • Suppression • SRI-HD • Edge enhance	• Gain • Frame averaging • Frequency • Scanning size (FOV or angle – depending on the probe, see probe specifications)
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Digital M-Mode

Adjustable	• Acoustic power • Dynamic range • Frequency • M colorization • Rejection	• Gain • Gray scale map • Sweep speed • M display format
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Anatomical M-Mode

M-Mode cursor adjustable at any plane

Can be activated from a Cine loop from a live or stored image

M and A capability

Available with Color Flow Mode

Curved Anatomical M-Mode

General System Parameters *(cont.)*

Digital Spectral Doppler Mode

Adjustable	• Acoustic power • Dynamic range • Transmit frequency • PW colorization • Sweep speed • Sample volume length • Spectrum inversion • Baseline shift • Time resolution • Compression • Trace sensitivity	• Gain • Gray scale map • Wall filter • Velocity scale range • Angle correction • Steered linear • Trace method • Doppler auto trace • Trace direction
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Digital Color Flow Mode

Adjustable	• Acoustic power • Gain • Velocity scale range • Wall filter • Packet size • Spatial filter • Baseline shift • Threshold • Sample volume control • Flash suppression	• Color maps, including velocity-variance maps • Line density • Steering angle • Frame average • Accumulation mode • Quantification (option)
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Digital Power Doppler Imaging

Adjustable	• Acoustic power • Gain • Velocity scale range • Wall filter • Packet size • Spatial filter • Frame average • Accumulation mode • Flash suppression	• Color maps including velocity-variance maps • Line density • Steering angle • Threshold • Sample volume control
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Continuous Wave Doppler (Option)

Adjustable	• Acoustic power • Dynamic range • Transmit frequency • CW colorization • Sweep speed • Angle correction • Trace method • Baseline shift • Compression • Trace direction	• Gain • Gray scale map • Wall filter • Velocity scale range • Spectrum inversion • Doppler auto trace • Trace sensitivity
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Available on the following probes: 3Sc-RS, 6S-RS, 12S-RS, 6Tc-RS, P2D, P6D and P8D

Automatic Optimization

Optimize B-Mode image to improve contrast resolution

Selectable amount of contrast resolution improvement (low, medium, high)

Auto TGC

Auto-spectral optimize adjusts	• Baseline • PRF (on live image)	• Invert • Angle correction
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Coded Harmonic Imaging

Available on all 2D and 4D probes

B-Flow/B-Flow Color (Option)

Available on C1-5-RS, 8C-RS, L6-12-RS, 12L-RS, 9L-RS, ML6-15-RS, L8-18i-RS, L4-12t-RS, L10-22-RS, L3-9i-RS, L12n-RS, E8CS-RS, BE9CS-RS, L3-12-RS and IC59-RS probes

Background: on/off

Sensitivity/PRI

Line density

Edge enhance

Frame average

Gray scale map

Tint map

Dynamic range

Rejection

Gain

Hybrid B-Flow	• Supported on C1-5-RS, 12L-RS, 9L-RS, ML6-15-RS, L4-12t-RS, L3-12-RS and L12n-RS • B & B-Flow simultaneous dual display • B & B-Flow overlay display
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B-Flow Color (BFC)

B-Flow High Definition Color (HD Color)	Supported on C1-5-RS, 12L-RS, ML6-15-RS, L4-12t-RS, L3-12-RS and L12n-RS
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Accumulation

Coded Contrast Imaging (Option)

Available on C1-5-RS, 9L-RS, 3Sc-RS, IC9-RS and BE9CS-RS probes

2 contrast timers

Timed updates: 0.05 – 10 seconds

Accumulation mode, six levels

Maximum Enhance Mode

General System Parameters *(cont.)*

Coded Contrast Imaging (Option) *(cont.)*

Flash

Time Intensity Curve (TIC) Analysis

Auto MI control

The LOGIQ P9 is designed for compatibility with commercially available ultrasound contrast agents. Because the availability of these agents is subject to government regulation and approval, product features intended for use with these agents may not be commercially marketed nor made available before the contrast agent is cleared for use. Contrast related product features are enabled only on systems for delivery to an authorized country or region of use.

LOGIQ View (Option)

Extended Field of View imaging

Available on the following probes: C1-5-RS, 8C-RS, L6-12-RS, 12L-RS, 9L-RS, ML6-15-RS, L8-18i-RS, L4-12t-RS, L10-22-RS, L3-9i-RS, L12n-RS, E8C-RS, E8CS-RS, IC9-RS, BE9CS-RS, RIC5-9A, 6Tc-RS, RAB2-6-RS, 3SC-RS, 6S-RS, 12S-RS

For use in B-Mode

CrossXBeam is available on linear probes

Auto detection of scan direction

Pre or post-process zoom

Rotation

Auto fit on monitor

Measurements in B-Mode

3D

Allows unlimited rotation and planar translations

3D reconstruction from Cine sweep

Advanced 3D (Option)

Acquisition of color data

Automatic rendering

3D landscape technology

3D movie

Real-time 4D (Option)

Acquisition modes	• Real-time 4D • Static 3D
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Real-time 4D (Option) *(cont.)*

Visualization modes	• 3D rendering (diverse surface and intensity projection modes) • Sectional planes (three section planes perpendicular to each other) • Volume contrast imaging-static (option) • Tomographic ultrasound imaging (option)
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Render mode	Surface texture, surface smooth, max-, min- and X-ray (average intensity projection), mix mode of two render modes
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Curved 3 point render start

3D movie

Scalpel: 3D cut tool

Display format	• Quad: A-/B-/C-Plane/3D • Dual: A-Plane/3D • Single: 3D or A- or B- or C-Plane
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Automated Volume Calculation – VOCAL II (option)

Betaview

Auto sweep

STIC (option)

HDlive (option)

Omniview (option)	VCI OmniView
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Scan Assistant (Option)

Factory programs

User defined programs

Steps include image annotations, mode transitions, basic imaging controls and measurement initiation

Shear Wave Elastography (Option)

Available on the following probes: C1-5-RS, L3-12-RS

User programmable measurement display in kPa and meters per sec.

Single and dual view display

B Steer+ (Option)

Available on the following probes: C1-5-RS, 8C-RS, L6-12-RS, 12L-RS, 9L-RS, ML6-15-RS, L4-12t-RS, L3-12-RS, L12n-RS, RAB2-6-RS

General System Parameters *(cont.)*

Strain Elastography (Option)

Available on C1-5-RS, L6-12-RS, 12L-RS, 9L-RS, ML6-15-RS, L4-12t-RS, L12n-RS, E8CS-RS, BE9CS-RS, L3-12-RS and IC9-RS probes

Semi-Quantification²

TVI (Option)

Myocardial doppler imaging with color overlay on tissue image

Available on the sector probes

Tissue color overlay can be removed to show just the 2D image, still retaining the tissue velocity information

Curved anatomical M-Mode: free (curved) drawing of M-Mode generated from the cursor independent from the axial plane

Q-Analysis: multiple time motion trace display from selected points in the myocardium

Stress Echo (Option)

Advanced and flexible Stress Echo examination capabilities

Provides exercise and pharmacological protocol templates

8 default templates

Template editor for user configuration of existing templates or creation of new templates

Reference scan display during acquisition for stress level comparison (dual screen)

Baseline level/previous level selectable

Raw data continuous capture

Over 100 sec. available

Wall motion scoring (bulls-eye and segmental)

Smart stress: automatically set up various scanning parameters (for instance, geometry, frequency, gain, etc.) according to same projection on previous level

Scan Assistant (Option)

Factory programs

User-defined programs

Steps include image annotations, mode transitions, basic imaging controls and measurement initiation

Compare Assistant (Option)

Allows side-by-side comparison of previous ultrasound and other modality exams during live scanning

Power Assistant (Option)

Allows moving the system without a complete system shutdown and boot-up power cycle

Extended battery for off line scanning (option) provides battery powered live scanning

Breast Productivity Package (Option)

Worksheet summary includes measurements and locations for nodule, parathyroid and lymph node

Feature assessment

BI-RADS® assessment

User editable

Thyroid Productivity Package (Option)

Worksheet summary includes measurements and locations for nodule, parathyroid and lymph node

Feature assessment

User editable

Auto EF (Option)

Allows semi-automatic measurement of the global EF (Ejection fraction)

User editable

Cardiac Strain (Cardiac AFI) (Option)

Allows assessing the left ventricle with all segments at a glance by combining three longitudinal views into one comprehensive bulls-eye view

2D strain based data moves into clinical practice

Virtual Convex

Provides a convex Field of View

Compatible with CrossXBeam

Available on all linear and sector transducers

SRI-HD

Speckle Reduction Imaging

Provides multiple levels of speckle reduction

Compatible with side-by-side DualView display

Compatible with all linear, convex and sector transducers

Compatible with B-Mode, color, contrast agent and 3D imaging

General System Parameters *(cont.)*

CrossXBeam

Provides 3, 5, 7 or 9 angles of spatial compounding

Live side-by-side DualView display

Compatible with	• Color Mode • SRI-HD • Virtual convex	• PW • Coded harmonic imaging
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Available on C1-5-RS, 8C-RS, L6-12-RS, 12L-RS, 9L-RS, ML6-15-RS, L8-18i-RS, L4-12t-RS, L10-22-RS, L3-9i-RS, L12n-RS, E8C-RS, E8CS-RS, BE9CS-RS, RIC5-9A-RS, IC9-RS, L3-12-RS and RAB2-6-RS probes

Controls Available While “Live”

Write zoom

B/M/CrossXBeam Mode

Gain

TGC

Dynamic range

Acoustic output

Transmission focus position

Transmission focus number

Line density control

Sweep speed for M-Mode

Number of angles for CrossXBeam

PW-Mode

Gain

Dynamic range

Acoustic output

Transmission frequency

PRF

Wall filter

Spectral averaging

Sample volume gate	• Length • Depth
--------------------	--

Velocity scale

Color Flow Mode

CFM gain

CFM velocity range

Acoustic output

Controls Available While “Live” *(cont.)*

Wall echo filter

Packet size

Frame rate control

CFM spatial filter

CFM frame averaging

CFM line resolution

Frequency/velocity baseline shift

Controls Available on “Freeze” or Recall

Automatic optimization

SRI-HD

CrossXBeam – display non-compounded and compounded image simultaneously in split screen

3D reconstruction from a stored Cine loop

B/M/CrossXBeam Mode

Gray map optimization

TGC

Colorized B and M

Frame average (loops only)

Dynamic range: Anatomical M-Mode

Max read zoom to 8x: baseline shift

Sweep speed

PW Mode

Gray map

Post gain

Baseline shift

Sweep speed

Invert spectral wave form

Compression

Rejection

Colorized spectrum

Display format

Doppler audio

Angle correct

Quick angle correct

Auto angle correct

General System Parameters (cont.)

Controls Available on “Freeze” or Recall (cont.)
Color flow
Overall gain (loops and stills)
Color map
Transparency map
Frame averaging (loops only)
Flash suppression
CFM display threshold
Spectral invert for Color/Doppler
Anatomical M-Mode on Cine loop

Measurements/Calculations

General B-Mode
Depth and distance
Circumference (ellipse/trace)
Area (ellipse/trace)
Volume (ellipsoid)
% Stenosis (area or diameter)
Angle between two lines

General M-Mode
M-Depth
Distance
Time
Slope
Heart rate

General Doppler Measurements/Calculations
Velocity
Time
A/B ratio (velocities/frequency ratio)
PS (Peak Systole)
ED (End Diastole)
PS/ED (PS/ED ratio)
ED/PS (ED/PS ratio)
AT (Acceleration Time)

General Doppler Measurements/Calculations (cont.)
ACCEL (Acceleration)
TAMAX (Time Averaged Maximum Velocity)
Volume Flow (TAMEAN and vessel area)
Heart rate
PI (Pulsatility Index)
RI (Resistivity Index)

Real-time Doppler Auto Measurements/Calculations
PS (Peak Systole)
ED (End Diastole)
MD (Minimum Diastole)
PI (Pulsatility Index)
RI (Resistivity Index)
AT (Acceleration Time)
ACC (Acceleration)
PS/ED (PS/ED ratio)
ED/PS (ED/PS ratio)
HR (Heart Rate)
TAMAX (Time Averaged Maximum Velocity)
PVAL (Peak Velocity Value)
Volume Flow (TAMEAN and vessel area)

OB Measurements/Calculations	
Gestational age by	• GS (Gestational Sac) • CRL (Crown Rump Length) • FL (Femur Length) • BPD (Biparietal Diameter) • AC (Abdominal Circumference) • HC (Head Circumference) • APTD x TTD (Anterior/Posterior Trunk Diameter by Transverse Trunk Diameter) • FTA (Fetal Trunk cross-sectional Area) • BD (Binocular Distance) • HL (Humerus Length) • FT (Foot Length) • OFD (Occipital Frontal Diameter) • TAD (Transverse Abdominal Diameter) • TCD (Transverse Cerebellum Diameter) • THD (Thorax Transverse Diameter) • TIB (Tibia Length) • ULNA (Ulna Length)

Measurements/Calculations *(cont.)*

OB Measurements/Calculations *(cont.)*

Estimated fetal weight (EFW) by	• AC, BPD • AC, BPD, FL, HC • AC, FL, HC • BPD, APTD, TTD, FL	• AC, BPD, FL • AC, FL • AC, HC • BPD, APTD, TTD, SL
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Calculations and ratios	• FL/BPD • FL/AC • FL/HC • HC/AC • CI (Cephalic Index) • AFI (Amniotic Fluid Index) • CTAR (Cardio-Thoracic Area Ratio) • MCA PS (Middle Cerebral Artery Peak Systolic Velocity) • MCA CP (Middle Cerebral Artery Pulsatility Index Over Umbilical Artery Pulsatility Index Ratio) • MCA PI (Middle Cerebral PI) • MCA RI (Middle Cerebral RI) • UmbArt PI (Umbilical artery PI) • UmbArt RI (Umbilical artery RI) • UtArt PI (Uterine artery PI) • UtArt RI (Uterine artery RI)
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Measurements/calculations by: ASUM, ASUM 2001, Berkowitz, Bertagnoli, Brenner, Campbell, CFEF, Chitty, Eik-Nes, Ericksen, Goldstein, Hadlock, Hansmann, Hellman, Hill, Hohler, Jeanty, JSUM, Kurtz, Mayden, Mercer, Merz, Moore, Nelson, Osaka University, Paris, Rempen, Robinson, Shepard, Shepard/Warsoff, Tokyo University, Tokyo/Shinozuka, Yarkoni

Fetal graphical trending

Growth percentiles

Multi-gestational calculations (4)

Fetal qualitative description (anatomical survey)

Fetal environmental description (biophysical profile)

Programmable OB tables

Over 20 selectable OB calculations

Expanded worksheets

GYN Measurements/Calculations

Right ovary length, width, height

Left ovary length, width, height

Uterus length, width, height

Cervix length, trace

Ovarian volume

ENDO (Endometrial Thickness)

Ovarian RI

GYN Measurements/Calculations *(cont.)*

Uterine RI

Follicular measurements

Summary reports

IOTA (International Ovarian Tumor Analysis) LR2 worksheet

Vascular Measurements/Calculations

SYS DCCA (Systolic Distal Common Carotid Artery)

DIAS DCCA (Diastolic Distal Common Carotid Artery)

SYS MCCA (Systolic Mid Common Carotid Artery)

DIAS MCCA (Diastolic Mid Common Carotid Artery)

SYS PCCA (Systolic Proximal Common Carotid Artery)

DIAS PCCA (Diastolic Proximal Common Carotid Artery)

SYS DICA (Systolic Distal Internal Carotid Artery)

DIAS DICA (Systolic Distal Internal Carotid Artery)

SYS MICA (Systolic Mid Internal Carotid Artery)

DIAS MICA (Diastolic Mid Internal Carotid Artery)

SYS PICA (Systolic Proximal Internal Carotid Artery)

DIAS PICA (Diastolic Proximal Internal Carotid Artery)

SYS DECA (Systolic Distal External Carotid Artery)

DIAS DECA (Diastolic Distal External Carotid Artery)

SYS PECA (Systolic Proximal External Carotid Artery)

DIAS PECA (Diastolic Proximal External Carotid Artery)

VERT (Systolic Vertebral Velocity)

SUBCLAV (Systolic Subclavian Velocity)

Automatic IMT

Summary Reports

Urological Calculations

Bladder volume

Prostate volume

Left/right renal volume

Generic volume

Post-void bladder volume

Probes

LOGIQ P9

C1-5-RS, 8C-RS, E8C-RS, E8CS-RS, BE9CS-RS, 9L-RS, 12L-RS, L8-18i-RS, L6-12-RS, L4-12t-RS, L10-22-RS, L3-9i-RS, ML6-15-RS, L12n-RS, 3Sc-RS, 6S-RS, 12S-RS, RAB2-6-RS, RIC5-9A-RS, P6D, P8D, L3-12-RS, IC9-RS, 6Tc-RS, P2D

C1-5-RS

Convex probe

Applications	Abdomen, Vascular, OB/GYN, Urology
Biopsy guide	Multi-angle, disposable with a reusable bracket (H40432LE)

8C-RS

Micro convex probe

Applications	Neonatal, Pediatrics
Biopsy guide	No

E8C-RS

Endocavitary micro convex probe

Applications	OB/GYN, Urology, Transvaginal, Transrectal
Biopsy guide	Single-angle, disposable with a disposable bracket (E8385MJ, E8333JB), single-angle, reusable bracket (H40412LN)

E8CS-RS

Endocavitary micro convex probe

Applications	OB/GYN, Urology, Transvaginal, Transrectal
Biopsy guide	Single-angle, disposable with a disposable bracket (E8385MJ, E8333JB), single-angle, reusable bracket (H40412LN)

IC9-RS

Endocavitary micro convex probe

Applications	OB/GYN, Urology, Transvaginal, Transrectal
Biopsy Guide	Single-angle, disposable with a disposable bracket (H48691YW), single-angle, reusable bracket (H48701MN)

BE9CS-RS

Endocavitary micro convex probe

Applications	Urology, Transrectal
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BE9CS-RS (cont.)

Biopsy guide	Single-angle, disposable with a disposable bracket (E8387M, H42742LH, H42742LJ), single-angle, reusable bracket (E8387MA)
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RAB2-6-RS

Convex volume probe

Applications	Abdomen, OB/GYN, Urology
Biopsy guide	Multi-angle, disposable with reusable bracket (H48681ML)

RIC5-9A-RS

Endocavitary micro convex volume probe

Applications	OB/GYN, Urology, Transvaginal, Transrectal
Biopsy guide	Single-angle, disposable with a disposable bracket (H48681GF), single-angle, reusable bracket (H46721R)

9L-RS

Linear probe

Applications	Vascular, Small Parts, Pediatric, Abdomen
Biopsy guide	Multi-angle, disposable with a reusable bracket (H4906BK)

12L-RS

Linear probe

Applications	Vascular, Small Parts, Neonatal, Pediatrics, Musculoskeletal
Biopsy guide	Multi-angle, disposable with a reusable bracket (H40432LC)

L8-18i-RS

Linear probe

Applications	Vascular, Small Parts, Neonatal, Pediatrics, Intraoperative
Biopsy guide	No

L6-12-RS

Linear probe

Applications	Abdomen, Vascular, Small Parts, Pediatrics, Neonatal, Musculoskeletal
Biopsy guide	Multi-angle, disposable with a reusable bracket (H40432LC)

Probes *(cont.)*

L12n-RS

Linear probe

Applications	Interventional Guidance, Vascular, Small Parts, Neonatal, Pediatrics, Musculoskeletal
Biopsy guide	Multi-angle, disposable with a reusable bracket. Infinite-angle (in plane biopsy kit), disposable with a reusable bracket. 4 configurable buttons to support various operation.

L4-12t-RS

Linear probe

Applications	Small Parts, Vascular, Pediatrics, Neonatal, Musculoskeletal
Biopsy guide	Multi-angle, disposable with a reusable bracket (H40432LC). Single-angle, disposable with a reusable bracket (H48392LT: free hand, H48392LL: transverse)

L10-22-RS

Linear probe

Applications	Small Parts, Musculoskeletal, Neonatal
Biopsy guide	N/A

L3-9i-RS

Linear probe

Applications	Small Parts, Vascular, Musculoskeletal, Intraoperative
Biopsy guide	N/A

ML6-15-RS

Matrix array linear probe

Applications	Small Parts, Vascular, Neonatal, Pediatrics, Musculoskeletal
Biopsy guide	Multi-angle, disposable with a reusable bracket (H40432LJ)

L3-12-RS

Linear Probe

Applications	Vascular, Small Parts, Neonatal, Pediatrics, Abdomen
Biopsy Guide	Multi-angle, disposable with a reusable bracket (H48302AA)

3Sc-RS

Phased array sector probe

Applications	Cardiac, Transcranial, Abdomen
Biopsy guide	Multi-angle, disposable with a reusable bracket (H46222LC)

6S-RS

Phased array sector probe

Applications	Cardiac Neonatal, Pediatrics
Biopsy guide	No

12S-RS

Phased array sector probe

Applications	Pediatrics, Neonatal
Biopsy guide	N/A

6Tc-RS

TEE Sector (Trans-esophageal) Probe

Applications	Cardiac (Transesophageal)
Biopsy Guide	None

P6D

CW split crystal probe

Applications	Cardiac, Vascular
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P8D

CW split crystal probe

Applications	Cardiac, Vascular
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P2D

CW Split Crystal Probe

Applications	Cardiac, Vascular
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Inputs and Outputs

HDMI out

Ethernet network (RJ45)

S-video out

Composite video out

USB (2x in front (USB 3.0), 3x in rear, 2x monitor)

AC power input

Pinpoint™ GT Needle Guidance Technology

Needle Guidance Technology – Optional

Accurate magnetic needle tracking (± 1.45 mm)

Pinpoint™ GT Needle Guidance Technology practice kit

Application with Pinpoint™ GT Needle Guidance Technology	Anesthesia, Musculoskeletal, Nerve Block
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Flexible needle selection	• From list manually • From label with Barcode Reader automatically
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Comprehensive multi-view	• Front View • Side View • Top View
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Safety Conformance

Classified to ANSI/AAMI ES60601-1 2005 R1 2012 Medical Electrical Equipment, Part 1: General Requirements for Safety by a Nationally Recognized Test Lab

Certified to CSA CAN/CSA-C22.2 NO. 60601-1:14 General requirements for safety

CE Marked to Council Directive 93/42/EEC on Medical Devices
Conforms to the following standards for safety:

IEC/EN 60601-1 2nd Edition Medical electrical equipment – Part 1: General requirements for safety

IEC/EN 60601-1 3.1 Edition. Medical electrical equipment – Part 1: General requirements for basic safety and essential performance

IEC/EN 60601-1-1 Medical electrical equipment – Part 1-1: General requirements for safety – Collateral Standard: Safety requirements for medical electrical systems

IEC/EN 60601-1-2 Medical electrical equipment – Part 1-2: General requirements for safety – Collateral Standard: Electromagnetic compatibility – requirements and tests

IEC/EN 60601-1-4 Medical electrical equipment Part 1- 4: General requirements for safety – Collateral Standard: programmable electrical medical systems

IEC/EN 60601-1-6 Medical electrical equipment Part 1 -6: General requirements for basic safety and essential performance – Collateral Standard: Usability

IEC/EN 60601-2-18 Medical electrical equipment – Part 2-18: Particular requirements for the basic safety and essential performance of endoscopic equipment

IEC/EN 60601-2-37 Medical electrical equipment – Part 2-37: Particular requirements for the safety of ultrasonic medical diagnostic and monitoring equipment

IEC/EN 62366 Application of usability engineering to medical devices

IEC/EN 62304 Software Life Cycle Processes

IEC/EN 62359 Ultrasonic – Field characterization – Test methods for the determination of thermal and mechanical indices related to medical diagnostic ultrasonic fields

EN ISO 15223-1: Symbols to be used with medical device labels, labelling and information to be supplied

ISO 10993-1 Biological evaluation of medical devices – Part 1 Evaluation and testing

NEMA UD2 Acoustic output measurement standard for diagnostic ultrasound equipment

NEMA UD3 Standard for real time display of thermal and mechanical acoustic output indices on diagnostic ultrasound equipment (MI, TIS, TIB, TIC)

EMC Emissions Group 1, Class B device requirements as per Sub clause 4.2 of CISPR 11

WEEE (Waste Electrical and Electronic Equipment)

ROHS according to 2011/65/EU Including national deviations

1. The LOGIQ P9 is a highly mobile and easy to use, performance multi-purpose color Doppler imaging system, designed for Abdominal, Small Parts, Musculoskeletal, Breast, Vascular, Cardiology, Transcranial, Urology, Pediatric, Neonatal, Obstetrics and Gynecology applications.
2. Contrast Enhanced Ultrasound is available in the U.S. for characterization of focal liver lesions and left ventricle opacity only.
3. Elastography with semi-Quantification (Elastography Quantification) described in this material has not been cleared by the U.S. FDA and is not available for promotion or sale in the United States.

Imagination at work

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LOGIQ™ P9/P7

Probe Guide



The LOGIQ P9/P7 is a highly capable ultrasound system that provides excellent image quality and productivity through easy to use tools across a wide range of applications in a portable, ergonomic, budget-friendly system design.

	Description	Applications	FOV	Bandwidth	Biopsy Guide	System
Convex Array						
 C1-5-RS H40462LA	Wideband convex array probe	Abdomen, OB/GYN, Urology, Vascular	70°	1 – 6 MHz	Multi-angle, disposable with a reusable bracket (H40432LE)	LOGIQ P9
 4C-RS H40402LS	Wideband convex array probe	Abdomen, OB/GYN, Urology, Vascular	58°	1 – 5 MHz	Multi-angle, disposable with a reusable bracket (E8385NA)	LOGIQ P7
Micro-convex Array						
 8C-RS H40402LS	Wideband micro-convex array probe	Neonatal, Pediatrics	132°	3 – 11 MHz	No	LOGIQ P9 LOGIQ P7
 E8C-RS H40402LN	Wideband micro-convex intra-cavitary array probe	OB/GYN, Urology, Endocavity	132°	3 – 11 MHz	Single-angle, disposable with a disposable bracket (E8385MJ, E8333JB), Single-angle, reusable bracket (H40412LN)	LOGIQ P9 LOGIQ P7
 E8Cs-RS H48062AF	Wideband micro-convex intra-cavitary array probe	OB/GYN, Urology, Endocavity	168°	3 – 11 MHz	Single-angle, disposable with a disposable bracket (E8385MJ, E8333JB), single-angle, reusable bracket (H40412LN)	LOGIQ P9 LOGIQ P7
 IC9-RS H48691PJ	Wideband micro-convex intracavity array probe	OB/GYN, Urology, Endocavity	168°	2 – 11 MHz	Single-angle, disposable with a disposable bracket (H48691YW), Single-angle, reusable bracket (H48701MN)	LOGIQ P9 LOGIQ P7
 BE9CS-RS H40482LN	Wideband micro-convex intra-cavitary bi-plane array probe	Urology, Transrectal	127° x 2	3 – 12 MHz	Single-angle, reusable (E8387MA), disposable (E8387M), disposable starter kit (H42742LH), disposable starter kit (H42742LJ)	LOGIQ P9 LOGIQ P7
Linear Array						
 ML6-15-RS H40462LM	Wideband linear matrix array probe	Small Parts, Vascular, Neonatal, Pediatrics, Musculoskeletal	50 mm	4 – 15 MHz	Multi-angle, disposable with a reusable bracket (H40432LJ)	LOGIQ P9
 L3-12-RS H44901AP	Wideband Linear Array Probe	Vascular, Small Parts, Neonatal, Pediatrics, Abdomen	51.2 mm	2 – 11 MHz	Multi-angle, disposable with a reusable bracket (H48032AA)	LOGIQ P7 LOGIQ P9
 L10-22-RS H48312AH	Wideband linear array probe	Small Parts, Neonatal, Musculoskeletal	13 mm	7 – 20 MHz	No	LOGIQ P9

	Description	Applications	FOV	Bandwidth	Biopsy Guide	System
Linear Array (cont.)						
 12L-RS H40402LY	Wideband linear array probe	Small Parts, Vascular, Pediatrics, Neonatal, Musculoskeletal	38 mm	3 – 12 MHz	Multi-angle, disposable with a reusable bracket (H40432LC)	LOGIQ P9
 L12n-RS H48062AH	Wideband linear array probe	Interventional Guidance, Vascular, Small Parts, Neonatal, Pediatrics, Musculoskeletal	38 mm	3 – 12 Mhz	Multi-angle, disposable with a reusable bracket (H40432LC), multi-angle, disposable with a reusable bracket (H48392LL), multi-angle, disposable with a reusable bracket (H48392LT)	LOGIQ P9 LOGIQ P7
 L4-12t-RS H48062AB	Wideband linear array probe	Small Parts, Vascular, Pediatric, Neonatal, Musculoskeletal	38 mm	3 – 12 MHz	Multi-angle, disposable with a reusable bracket (H40432LC), multi-angle, disposable with a reusable bracket (H48392LL), multi-angle, disposable with a reusable bracket (H48392LT)	LOGIQ P9 LOGIQ P7
 9L-RS H40442LL	Wideband linear array probe	Vascular, Small Parts, Pediatrics, Abdomen	44 mm	2 – 8 MHz	Multi-angle, disposable with a reusable bracket (H4906BK)	LOGIQ P9 LOGIQ P7
 L6-12-RS H48062AC	Wideband linear array probe	Small Parts, Vascular, Pediatrics, Neonatal, Abdomen	38.4 mm	5 – 11 MHz	Multi-angle, disposable with a reusable bracket (H40432LC)	LOGIQ P9 LOGIQ P7
 L8-18i-RS H40462LF	Wideband linear array probe	Small Parts, Vascular, Pediatrics, Neonatal, Intraoperative	25 mm	4 – 15 MHz	No	LOGIQ P9
 L3-9i-RS H46442LK	Wideband linear array probe	Small Parts, Vascular, Musculoskeletal, Intraoperative	38 mm	2 – 9 MHz	No	LOGIQ P9
Sector Array						
 3Sc-RS H45041DL	Wideband sector array probe	Cardiac, Abdomen, Transcranial	120°	1 – 5 MHz	Multi-angle, disposable with a reusable bracket (H46222LC)	LOGIQ P9 LOGIQ P7
 6S-RS H45021RP	Wideband sector array probe	Cardiac, Neonatal, Pediatric	90°	2 – 8 MHz	No	LOGIQ P9 LOGIQ P7
 12S-RS H44901AB	Wideband sector array probe	Pediatric, Neonatal	90°	4 – 12 MHz	No	LOGIQ P9 LOGIQ P7

	Description	Applications	FOV	Bandwidth	Biopsy Guide	System
	Sector Array (cont.)					
	TEE probe	Cardiac	90°	2 – 8 MHz	No	LOGIQ P9
	Real-time 4D					
	Wideband real-time 4D probe	Abdomen, OB/GYN, Urology	66° (B), 85° (Volume scan)	1 – 5 MHz	Multi-angle, disposable with a reusable bracket (H48681ML)	LOGIQ P9 LOGIQ P7
	Wideband real-time 4D intra-cavitary probe	Endocavity, OB/GYN, Urology	146° (B) 120° (Volume angle)	3 – 10 MHz	Single-angle, reusable bracket (H46721R), Single-angle, disposable (H48681GF)	LOGIQ P9 LOGIQ P7
	Specialty					
	CW split crystal pencil probe	Cardiac, Vascular	N/A	8 MHz	No	LOGIQ P9 LOGIQ P7
	CW split crystal pencil probe	Cardiac, Vascular	N/A	6 MHz	No	LOGIQ P9 LOGIQ P7
	CW split crystal pencil probe	Cardiac, Vascular		2 MHz	No	LOGIQ P9 LOGIQ P7

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Product may not be available in all countries and regions. Full product technical specification is available upon request. Contact a GE Healthcare Representative for more information. Please visit www.gehealthcare.com/promotional-locations. Data subject to change.

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