

This is a glossary of symbols on Hologic® Breast and Skeletal Health products and labels. Use this glossary as a supplement to your product manuals.

Symbol	Description	Standard
	Prescription use only	FDA 21 CFR 801.109
	Not safe for magnetic resonance imaging	ASTM F2503 - 23e1, Reference 6.4.10.1; Fig. 10
	Conditional use for magnetic resonance imaging	ASTM F2503 - 23e1, Reference 6.4.6.1; Fig 6
	Safe for magnetic resonance imaging	ASTM F2503 - 23e1, Reference 6.4.4; Fig. 4
	Conditional use for magnetic resonance imaging implant and deployment system	Hologic per ASTM F2503 Reference 7.4.7
	Caution: Radiation	Hologic
	Warning: Radiation (X-rays produced)	Health Canada Radiation Emitting Device Regulations (CRC, c. 1370)
	Emergency stop	Hologic
	Provision for the disabled or handicapped persons	ISO 7000:2004, Reference 0100
	Footswitch connection	Hologic
	Footswitch cord management plate	Hologic
	Wi-Fi	Hologic

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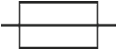
BSH Glossary of Symbols

Symbol	Description	Standard
	Danger: Pinch Point	Hologic
	Bluetooth	Hologic
	Battery	Hologic
YYYY-MM-DD	Date format: YYYY represents the year MM represents the month DD represents the day	Hologic
	Circuit breaker	Hologic
Translations in Box	Translations in Box	Hologic
Temp Logger Activated	Temperature Logger activated	Hologic
Temp Logger SN	Temperature Logger Serial Number	Hologic
PN	Part Number	Hologic
REV	Revision	Hologic
Model	Model Number	Hologic
QTY	Quantity	Hologic
	Packaging Unit	ISO 7000 Reference No. 2794
SFW SHIPPED	Software version shipped	Hologic
SFW Version	Software version number	Hologic

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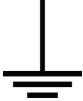
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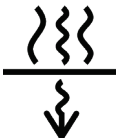
Symbol	Description	Standard
APPLICATION INSTALLATION	Application Installation	Hologic
Verified By	Verified By	Hologic
CASE	Case (box or package)	Hologic
VC	Vendor Code	Hologic
Made in USA	Made in United States of America	Hologic
Made in Costa Rica	Made in Costa Rica	Hologic
Assembled in Costa Rica	Assembled in Costa Rica	Hologic
Made in China	Made in China	Hologic
Made in Mexico	Made in Mexico	Hologic
Made in Thailand	Made in Thailand	Hologic
Patents	Patents	Hologic
USA REP	USA Representative	Hologic
LATERAL ARM USE ONLY	Lateral Arm Use Only	Hologic
Q.C. Use Only	Quality Control Use only	Hologic
UNLOAD THIS SIDE	Unload this side	Hologic
RAMP THIS SIDE	Ramp this side	Hologic
MATL	Material	Hologic

Symbol	Description	Standard
	Unlocking	IEC 60417, Reference 5570
	"ON" (power)	IEC 60417, Reference 5007
	"OFF" (power)	IEC 60417, Reference 5008
	Fuse	IEC 60417, Reference 5016
	Protective earth (ground)	IEC 60417, Reference 5019
	Equipotentiality	IEC 60417, Reference 5021
	Alternating Current	IEC 60417, Reference 5032
	Three-Phase Alternating Current with Neutral Conductor	IEC 60417, Reference 5032-2
	Dangerous voltage	IEC 60417, Reference 5036
	Non-ionizing electromagnetic radiation	IEC 60417, Reference 5140
	Type BF applied part	IEC 60417, Reference 5333
	Type B applied part	IEC 60417, Reference 5840
	Computer network	IEC 60417, Reference 5988
	Warning: Electricity	ISO-7010:2019, Reference W012

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Symbol	Description	Standard
	Caution: Risk of electric shock	IEC 60417, Reference 6042
	Note	Hologic
	USB	Hologic
	Universal Serial Bus (USB), port/plug	ISO 7000:2019, Reference 3650
	Locking: General	IEC 60417, Reference 5569
	Stand-by	IEC 60417, Reference 5009
	"ON"/"OFF" (push-push)	IEC 60417, Reference 5010
	Earth (ground)	IEC 60417, Reference 5017
	Electrostatic sensitive devices	IEC 60417, Reference 5134
	"ON" for a part of the equipment	IEC 60417, Reference 5264
	"OFF" for a part of the equipment	IEC 60417, Reference 5265
	Stand-by or preparatory state for a part of equipment	IEC 60417, Reference 5266
	X-ray source assembly	IEC 60417, Reference 5338

Symbol	Description	Standard
	X-ray source emitting	IEC 60417, Reference 5339
	Radiation filter	IEC 60417, Reference 5381
	Do not immerse in any liquid	IEC 60417, Reference 5995
IPX8	The equipment of accessory is suitable for continuous immersion in water (up to 1m of submersion for 1 hour)	IEC 60529
 IPX4	Splash-proof	IEC 60529
IPX6	Water projected in powerful jets against the equipment or accessory from any direction shall have no harmful effects	IEC 60529
IP54	Protected against dust limited ingress, no harmful deposits and protected against water splashed from all directions, limited ingress permitted	IEC 60529
IP22	Protected against solid foreign objects of 12,5 mm Ø and greater And Protection against vertically falling water drops when ENCLOSURE tilted up to 15°	IEC 60529
	Medical device	ISO 15223-1:2021, Reference 5.7.7
	Authorized Representative in the European Community	ISO 15223-1:2021, Reference 5.1.2
	Authorized Representative in the European Community	ISO 15223-1:2021/Amd 1:2025, Reference 5.1.2
	Indicates the authorized representative in Switzerland	MedDO; SR 812.213

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Symbol	Description	Standard
	Authorized Representative in the identified country of jurisdiction	ISO 15223-1:2021/Amd 1:2025, Reference 5.1.2
	Manufacturer	ISO 15223-1:2021, Reference 5.1.1
	Date of manufacture	ISO 15223-1:2021, Reference 5.1.3
	Country and date of manufacture	ISO 15223-1:2021, Reference 5.1.1/5.1.3
	Use-by Date	ISO 15223-1:2021, Reference 5.1.4
	Batch code	ISO 15223-1:2021, Reference 5.1.5
	Catalog number	ISO 15223-1:2021, Reference 5.1.6
	Serial number	ISO 15223-1:2021, Reference 5.1.7
	Sterilized using ethylene oxide	ISO 15223-1:2021, Reference 5.2.3
	Sterilized using irradiation	ISO 15223-1:2021, Reference 5.2.4
	Single sterile barrier system	ISO 15223-1:2021 Reference 5.2.11 (ISO 7000-3707)
	Double sterile barrier system	ISO 15223-1:2021 Reference 5.2.12 (ISO 7000-3704)
	Single sterile barrier system with protective packaging inside	ISO 15223-1:2021 Reference 5.2.13 (ISO 7000-3708)
	Single sterile barrier system with protective packaging outside	ISO 15223-1:2021 Reference 5.2.14 (ISO 7000-3709)

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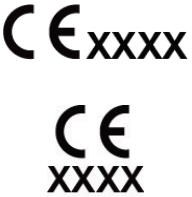
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Symbol	Description	Standard
	Do not re sterilize	ISO 15223-1:2021, Reference 5.2.6
	Non-sterile	ISO 15223-1:2021, Reference 5.2.7
	Do not use if package is damaged and consult instructions for use OR Do not use if the product sterile barrier system or its packaging is compromised	ISO 15223-1:2021, Reference 5.2.8
	Fragile; handle with care	ISO 15223-1:2021, Reference 5.3.1
	Keep dry	ISO 15223-1:2021, Reference 5.3.4
	Lower limit of temperature	ISO 15223-1:2021, Reference 5.3.5
	Upper limit of temperature	ISO 15223-1:2021, Reference 5.3.6
	Temperature limit	ISO 15223-1:2021, Reference 5.3.7
	Humidity limitation	ISO 15223-1:2021, Reference 5.3.8
	Atmospheric pressure limitation	ISO 15223-1:2021, Reference 5.3.9
	Do not re-use	ISO 15223-1:2021, Reference 5.4.2
 www.hologic.com/package-inserts	URL Location of Hologic Instructions for Use	Hologic per ISO 15223-1:2021, Reference 5.4.3

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Symbol	Description	Standard
	Consult Instructions for Use; operating manual	ISO 15223-1:2021, Reference 5.4.3 (ISO 7000, Reference 1614)
	Electronic Instructions for Use	ISO 9687-1:2018 (ISO 7000, Reference 3500)
	Non-pyrogenic	ISO 15223-1:2021, Reference 5.6.3 (ISO 7000, Reference 2724)
	Refer to instruction manual/booklet; instruction manual/booklet must be read	ISO 7010:2019, Reference M002
	Caution	ISO 15223-1:2021, Reference 5.4.4 (ISO 7000, Reference 0434A)
	Mass; weight	ISO 7000:2004, Reference 1321A
	This way up	ISO 7000:2014, Reference 0623
	Do not stack	ISO 7000:2004, Reference 2402 IEC TR 60878:2022
	Stacking limit by number	ISO 7000:2014, Reference 2403 IEC TR 60878:2022
	Warning	ISO 7010:2019, Reference W001
	CE Mark European Conformity	MDR Regulation (EU) 2017/745

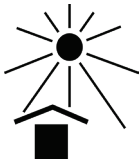
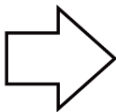
Symbol	Description	Standard
	CE Mark with Notified Body reference number	MDR Regulation (EU) 2017/745
	Contents	Hologic
	Contains or presence of natural rubber latex	ISO 15223-1:2021, Reference 5.4.5
	Does not contain natural rubber latex	Hologic
	Do not cut	Hologic
	CSA Group Certification Mark (Canada and US)	Recognized and accredited by the Standards Council of Canada (SCC) and the U.S. Occupational Safety and Health Administration (OSHA)
	General medical equipment as to electrical shock, fire, and mechanical hazards only in accordance with ANSI/AAMI ES 60601-1:2005 (AI:2012)	UL Classification
	General medical equipment as to electrical shock, fire, and mechanical hazards only in accordance with ANSI/AAMI ES 60601-1:2005 (AI:2012), CAN/CSA C22.2 No. 60601-1 (2014), and IEC 60601-1:2012	UL Classification (Canada and U.S.)

Symbol	Description	Standard
	ETL Listed Mark – Intertek (US)	IEC 60601-1 compliant, US
	ETL Listed Mark – Intertek (Canada and US)	IEC 60601-1 compliant, Canada and US
	INMETRO & UL-BR Mark (OCP 0029)	UL-BR INMETRO certification (Brazil)
	INMETRO & LMP-BR Mark (OCP 0131)	LMP-BR INMETRO certification (Brazil)
	INMETRO & LMP-BR Mark (OCP 0131)	LMP-BR INMETRO certification (Brazil)
	Recyclable symbol	Unicode, Reference U+267C
	Discard electrical and electronic equipment separately from standard waste. Send decommissioned material to Hologic or contact your service representative.	WEEE Directive 2012/19/EU
	Patient Identification	ISO 15223-1:2021, Reference 5.7.3
	Patient Information Website	ISO 15223-1:2021, Reference 5.7.4

Symbol	Description	Standard
	Health Care Centre or Doctor	ISO 15223-1:2021, Reference 5.7.5
	Date	ISO 15223-1:2021, Reference 5.7.6
	Unique Device Identifier	ISO 15223-1:2021, Reference 5.7.10
	Reject if Dot Black	Hologic
	No pushing	ISO 7010:2019, Reference P017
	No sitting	ISO 7010:2019, Reference P018
	No stepping on surface	ISO 7010:2019, Reference P019
	Ionized radiation emitted when x-ray beam is energized	ISO 7010:2019, Reference W003
	Warning: Biological hazard	ISO 7010:2019, Reference W009
	Lifting Hazard	Hologic

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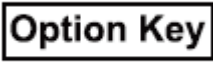
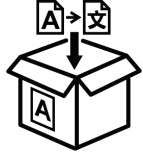
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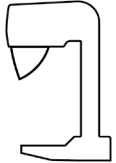
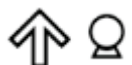
Symbol	Description	Standard
	Keep away from sunlight	ISO 15223-1:2021 Reference 5.3.2. (ISO 7000-0624)
	Contains or presence of phthalate; DEHP	BS EN 15986:2011, Reference A.4
	Contains or presence of phthalate; DEHP, BBP, and DBP	BS EN 15986:2011, Reference A.5
	Does not contain or have presence of phthalate	Hologic per BS EN 15986:2011, Reference A.4
	Contains hazardous substances	ISO 15223-1:2021 Reference 5.4.10 (ISO 7000-3723)
	Country code for translation	ISO 3166-1:2020
	Language code for translation	ISO 639:2023
	Warning: Moving Gears	ISO 7010:W089
	Functional Arrow	ISO 7000:2004, Reference 0251
	Warning: Laser Beam	ISO 7010:2019, Reference W004

Symbol	Description	Standard
	Warning: explosive material	ISO 7010:2019, Reference W002
	Warning: Non-ionizing radiation	ISO 7010:2019, Reference W005
	Warning: Floor level obstacle	ISO 7010:2019, Reference W007
	Warning: Toxic material	ISO 7010:2019, Reference W016
	Warning: Crushing	ISO 7010:2019, Reference W019
	Warning: Flammable materials	ISO 7010:2019, Reference W021
	Warning: Hot surface	ISO 7010:2019, Reference W017
	Caution: Hot surface	IEC 60417, Reference 5041
	Caution: Heavy End	Hologic
	Device driver and remote-control connection	Hologic
	Device driver cord management plate	Hologic

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Symbol	Description	Standard
	Saline tubing management plate	Hologic
	Option Key	Hologic
	Self-timer	IEC 60417, Reference 5879
	Call for maintenance	ISO-7000:2004, Reference 0717
	BioVision Camera	Hologic
	Apply specific torque	Hologic
	RCM	AS/NZS 4268 compliant for short-range radio equipment, Australian Standard AS/NZS 4417.1, and AS/NZS 4417.2.
	Translation	ISO 7000:2019, Reference 3728
	Translations in Box	Hologic per ISO 7000:2019, Reference 3728
	Compression – Footswitch	Hologic
	Compression Down	Hologic

Symbol	Description	Standard
	Compression Up	Hologic
	C-Arm	Hologic
	C-Arm Down	Hologic
	C-Arm Up	Hologic
	C-Arm Tilt Backward (-)	Hologic
	C-Arm Tilt Forward (+)	Hologic
	C-Arm Zero	Hologic
	Paddle Shift Controls	Hologic
	Tube Arm Smart Rotation	Hologic
	Tube Arm Counterclockwise Rotation	Hologic
	Tube Arm Clockwise Rotation	Hologic