

iCONNECT

User Guide



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Patents

Manufactured under one or more of the following United States patents:

Niobe

6,352,363; 6,975,197; 7,019,610; 7,161,453; 7,305,263; 7,313,429; 7,495,537;
7,772,950; 7,966,059

Genesis RMN

7,774,046; 7690,619

Navigant

7,516,416; 7,537,570; 7,540,288; 7,540,866; 7,543,239; 7,627,361;
7,630,752; 7,657,075; 7,708,696; 7,751,867; 7,756,308; 7,761,133;
7,769,428; 7,831,294; 7,853,306; 8,024,024; 8,192,374; 8,369,934;
8,721,655; 9,314,222

Manufactured under the following European patent:

Niobe: EP 1 488 431, issued in Germany, France, and Great Britain.

Navigant: EP 1 682 024, issued in Germany, France, and Great Britain.

Other patents issued and pending.

Trademarks

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- *Odyssey Cinema* is a trademark of Stereotaxis, Inc., registered in the European Community.
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- *Rhythmia* and *Maestro 4000* are trademarks of Boston Scientific.
- *Ampere* and *EnSite Precision* are trademarks of St. Jude Medical.
- *HAT 500* is a registered trademark of Osypka.
- *AcQMap* is a registered trademark of Acutus Medical.

All other brand names, product names, or trademarks found in this document are the property of their respective owners.

EMC Directive Statement

EMC Directive Compliance	This equipment was tested and found to conform to the Medical Directive 93/42/EEC for electromagnetic compatibility. Compliance with this Directive is based upon compliance with the following harmonized standards:
Emissions:	IEC 60601-1-2 ed 4.0 (2014-02) CISPR11 ed 6.1 (2016-06), Class A (Professional Healthcare Facility Environment)
Immunity:	IEC 60601-1-2 ed 4.0 (2014-02), IEC 61000-4-2 ed 4.0 (2008-12), IEC 61000-4-3 ed 3.2 (2010-04), IEC 61000-4-4 ed 3.0 (2012-04), IEC 61000-4-5 ed 3.0 (2014-05) + AMD:2017, IEC 61000-4-6 ed 4.0 (2014 COR2015), IEC 61000-4-8 ed 2.0 (2009-09), IEC 61000-4-11 ed 2.0 (2004-03) + AMD1:2017

When operating this equipment, verify that other devices installed near it conform to the applicable EMC standards for that device. This equipment is designed to be installed and operated in a Professional Healthcare Facility Environment.

Safety Standard Statement

Safety Standard Compliance This equipment was tested and found to conform to the following IEC 60601-1 Medical Electrical Equipment General Requirements for basic safety and essential performance test specifications:

Standard: CAN/CSA-C22.2 No. 60601-1:14
ANSI/AAMI ES60601-1:2005+A2 (R2012) +A1
IEC 60601-1: 2005 + CORR. 1:2006 + CORR. 2:2007 + AM1:2012
(or IEC 60601-1: 2012 reprint)

Related documents



Niobe User Guide

Genesis RMN User Guide

Celsius Catheter Cable (001-009071-1) IFU

Magnoflush Catheter Cable (001-009073-1) IFU

CARTO System Cable (001-009075-1) IFU

Maestro System Cable (001-009077-1) IFU

Stockert System Cable (001-009078-1) IFU

Ampere System Cable (001-009079-1) IFU

HAT500 System Cable (001-009080-1) IFU

Navistar Adapter System Cable (001-009081-1) IFU

Respective documentation for compatible devices and/or systems.

Operating conditions

Temperature: 15°C to 30°C

Humidity: 20% to 75%, non-condensing

Atmospheric pressure: 70 kPa to 106 kPa

Storage and transport conditions

Temperature: -10°C to 50°C

Relative humidity: 20% to 95%

Atmospheric pressure: 70 kPa to 106 kPa

Equipment information

iCONNECT Model no.: 001-009040-1

Classification: Class II Defibrillation-proof Type CF Applied Part equipment

Mode of operation: Continuous

Performance specifications:

Parameter	Specification
System Weight	1.0 kg
Maximum Power Draw	0.25 Amp @ 24 VDC

Catheter Cable Model no.:

Celsius: 001-009071-1

Magnoflush: 001-009073-1

System Cable Model no.:

CARTO: 001-009075-1

Maestro: 001-009077-1

Stockert: 001-009078-1

Ampere: 001-009079-1

HAT500: 001-009080-1

Navistar Adapter: 001-009081-1



WARNING: No modification of the iCONNECT System is allowed. No user-serviceable parts are inside iCONNECT. The user should not remove the cover or attempt to disassemble any portion of the iCONNECT System.

Notice to the User and/or Patient

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

Cover art

Cover art © 2020 Stereotaxis, Inc.

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1 Overview

1.1 Introduction

iCONNECT is used in conjunction with a Stereotaxis magnetic navigation system (MNS). The iCONNECT System provides an electronics hub allowing users to connect a combination of compatible catheters, mapping systems, and ablation generators. In addition, the iCONNECT System gathers and provides catheter impedance values to the *Navigant*® Workstation (NWS) System where cardiac wall contact is determined.

1.2 About These Instructions

The purpose of this guide is to provide the user of the iCONNECT System with instructions for operating the equipment and using the *Navigant* software starburst indicator.

The instructions provide information for installing and maintaining the iCONNECT System and for understanding the reporting indicators on the equipment as well as through the *Navigant* software.

1.3 Indications

The iCONNECT System is an accessory device that indicates when the tip of a compatible catheter is in contact with the heart wall when used in conjunction with a compatible Magnetic Navigation System.

1.4 Companion Systems

iCONNECT may be used with the following magnetically-compatible devices:

- NAVISTAR® RMT Catheter
- NAVISTAR® THERMOCOOL® RMT Catheter
- CELSIUS® RMT Catheter
- CELSIUS® THERMOCOOL® RMT Catheter
- MedFact MagnoFlush Catheter

iCONNECT may be used with the following ablation generators:

- Osypka HAT 500® Generator
- Ampere™ RF Generator
- Maestro 4000™ Cardiac Ablation System
- Biosense Webster Stockert 70 RF Generator

iCONNECT may be used with the following mapping systems:

- CARTO® 3
- Acutus AcQMap®




iCONNECT communicates with a Stereotaxis MNS which integrates with a compatible, digital fluoroscopy system to provide real-time guidance to the physician during an interventional procedure.

1.5 Required Training

The Stereotaxis representative provides training and associated materials for safe and effective use of iCONNECT's primary operating functions based on the intended use of the system.






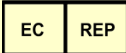



1.6 Graphics and Symbols

The following graphics and symbols are used in these instructions:

WARNING		WARNING indicates a potentially hazardous situation which, if not avoided, could result in death or serious injury.
CAUTION		CAUTION indicates a potentially hazardous situation which, if not avoided, could result in injury to patient or operator or damage to the equipment.
Note		Note identifies information that could affect the outcome or results of the procedure.

Warnings and Cautions precede the text and any procedure involving a clear risk to the operator(s), the patient, or the equipment. General Warnings and Precautions are listed earlier in the document in 1.6 Safety Instructions. Pay close attention to the instructions that accompany any warning, caution, and note symbols.

The following symbols are used in this document and/or on or near the system components:

Symbol	Name	Description
	Defibrillation-proof Type CF Applied Part	Indicates equipment suitable for direct cardiac application.
	Consult Instructions	See other material for additional information or instruction.
	Sterile	Sterile via ethylene oxide sterilization method.
	Non-Sterile	Device has not been sterilized.
	Manufacturer	Name and address of the manufacturer of the product.
	European Representative	Name and address of the authorized representative in the European Community.
	Reference	Part Number
	Lot Number	
	Expiration Date	

1.7 Glossary

The following terms appear in this document:

Term	Description
C-Arm	X-ray image intensifier that produces live X-ray images displayed on a monitor and named because of its configuration, with the top part of the C extending over the patient and the bottom extending under.
iCONNECT Electronics Hub	Hardware portion of the system that allows users to connect a combination of catheters, mapping systems, and ablation generators.
iCONNECT System	Gathers and provides catheter impedance values to <i>Navigant</i> where contact with the heart wall is determined.
EP	Electrophysiology.
Fluoro	Fluoroscopic, or Fluoroscopy.
Fluoroscope	X-ray image intensifier that produces live X-ray images displayed on a monitor. Also called X-ray or C-Arm.
Genesis Robotic Magnetic Navigation (RMN) System	A Stereotaxis system consisting of computer-controlled magnets that assist physicians in orienting and steering compatible, magnetically-adapted devices.
Impedance Graph	Feature of <i>Navigant</i> that helps the user determine cardiac wall contact.
Magnetic Navigation System (MNS)	A Stereotaxis system consisting of computer-controlled magnets that assist physicians in orienting and steering compatible, magnetically-adapted devices.
<i>Navigant</i> Workstation (NWS)	Platform of software applications designed to simplify clinical workflows. The <i>Navigant</i> product, when it does not include the <i>Odyssey</i> add-on, provides enhanced integration of catheterization and electrophysiology labs and improved automation during magnetic navigation of medical devices. (Available only in magnetic labs.)
Starburst	<i>Navigant</i> software feature of the iCONNECT System that indicates when the catheter tip is in contact with the heart wall.

1.8 Safety Instructions

1.8.1 Warnings

iCONNECT features do not replace the physician's knowledge, expertise, or judgment.



WARNING: US Federal law restricts this device to sale by or on the order of a physician.



WARNING: The iCONNECT System should be used only by qualified medical professionals who have been thoroughly trained in its use.



WARNING: The user should not attempt to upgrade, configure, or run any other software programs on the MNS computers, other than those specifically called out in the product documentation.



WARNING: The graphical representation of the catheter on the MNS display screen represents an approximate location and orientation of the catheter inside the patient's heart after the user acquires a new fluoro image and subsequently transfers it from the X-ray.



WARNING: The operator should always confirm device location using a live fluoroscopic image.



WARNING: Always verify catheter motion using live fluoroscopy images. Stop immediately if catheter motion can no longer be verified.



WARNING: If unexpected catheter motion occurs during ablation, stop RF energy delivery.



WARNING: If there is a cybersecurity breach during a procedure, press the Estop button to stop all device activities. Contact the Emergency call center to report the suspicious activity prior to resumption of the procedure.



WARNING: Catheter cables shall be re-sterilized after use to avoid potential for patient infection.



WARNING: Cables shall not be re-sterilized and used beyond indicated maximum as this could induce noise on ECG or mapping system.



WARNING: User shall not use devices for which packaging has been damaged.



WARNING: Non-approved companion cable systems shall not be connected as this could result in damage for either product.

1.8.2 Precautions



CAUTION: If a patient is to receive a permanent pacemaker or implantable cardiac defibrillator (ICD) *de novo*, the MNS magnets must be kept in the Stowed position. The magnets should remain in the Stowed position from the moment the ICD/pacemaker is brought inside the Procedure Room until the patient is removed from the Procedure Room. Proper functioning of ICD/pacemaker should be verified following the removal of the patient from the Procedure Room.



CAUTION: The points on the Mapping System screen are for reference only. Always confirm orientation of the catheter location with localization methods—fluoroscopy, for example—in addition to the electrograms.



CAUTION: The points and catheter tip location displayed on the Mapping System screen can be transferred to the *Navigant* screen. The catheter tip displayed on the *Navigant* screen is used only as a visual reference to aid navigation, not to identify the location of the catheter tip in the heart.



CAUTION: If the physician believes the catheter may be in the wrong location, the physician should refresh the fluoroscopy image. If a discrepancy occurs, the physician should reregister the mapping system to *Navigant*, remark the catheter base, and recalibrate the *Cardiodrive* System.

1.8.3 Notes



Always verify the fluoroscopy images match the current patient.



Before ablation, verify the field is applied to ensure proper device functionality.

1.9 Electromagnetic Compatibility Information



WARNING: The use of accessories, transducers, and cables other than those specified, with the exception of transducers and cables sold by Stereotaxis, Inc., may result in increased emissions or decreased immunity of the equipment.



WARNING: The equipment should not be used adjacent to or stacked with other equipment. If adjacent or stacked use is necessary, the equipment should be observed to verify normal operation in the configuration in which it will be used.



WARNING: This equipment has been tested for radiated radiofrequency (RF) immunity only at selected frequencies, and use of nearby emitters at other frequencies could result in improper operation. Improper operation may consist of (1) prevention of patient treatment, (2) uncontrolled motion of magnets, and (3) uncontrolled motion of the *Cardiodrive* System.



Note: In addition to the requirements of 7.9.3 of the general standard for Permanently Installed Large Medical Electrical (ME) Equipment and Large ME Systems for which the exemption specified in 8.6 from the testing requirements of IEC 61000-4-3 is used, this technical description includes the following information:

Following are the frequencies and modulations used to test the Immunity of the ME Equipment or ME System:

- 52.5Mhz
- 144Mhz
- 433Mhz
- 448Mhz
- 452Mhz
- 467Mhz
- 470Mhz
- 1.2Ghz
- 2.4Ghz
- 5.0Ghz

1.9.1 Emissions

The equipment is intended for use in the electromagnetic environment specified in the following tables. The customer or the user of the equipment should ensure that it is used in such an environment. The following table provides guidance and Stereotaxis’ declaration on **electromagnetic emissions** for the equipment:

Emissions	Compliance	Electromagnetic environment—guidance
RF emissions CISPR 11	Group 1	The equipment uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11	Class A Complies	
Harmonic emissions IEC 61000-3-2		
Voltage fluctuations/ flicker emissions IEC 61000-3-3		



WARNING: The EMISSIONS characteristics of this equipment make it suitable for use in industrial and hospital (CISPR 11 class A) settings. If it is used in a residential environment (for which CISPR 11 class B is normally required), this equipment might not offer adequate protection to radio-frequency communication services. The user might need to take mitigation measures, such as relocating or re-orienting the equipment.

1.9.2 Immunity—General electromagnetic effects


The following table provides guidance and Stereotaxis' declaration on **electromagnetic immunity** regarding general electromagnetic effects for the equipment:

Immunity test	IEC 60601 test level*	Compliance level*	Electromagnetic environment—guidance
Electrostatic discharge (ESD) IEC 61000-4-2	± 8 kV contact ± 2, 4, 8 & 15 kV air	± 8 kV contact ± 2, 4, 8 & 15 kV air	Floors should be wood, concrete, or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Electrical fast transient/burst IEC 61000-4-4	± 2 kV for power supply lines ± 1 kV for input/output lines	± 2 kV for power supply lines ± 1 kV for input/output lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	± 1 kV line(s) to line(s) ± 2 kV line(s) to earth	± 1 kV line(s) to line(s) ± 2 kV line(s) to earth	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	<5 % U_T (100 % dip in U_T) for 0.5 cycle 40 % U_T (60 % dip in U_T) for 5 cycles 70 % U_T (30 % dip in U_T) for 25 cycles <5 % U_T (>95 % dip in U_T) for 5 s	<5 % U_T (100 % dip in U_T) for 0.5 cycle 40 % U_T (60 % dip in U_T) for 5 cycles 70 % U_T (30 % dip in U_T) for 25 cycles <5 % U_T (>95 % dip in U_T) for 5 s	Mains power quality should be that of a typical commercial or hospital environment. If the user of the equipment requires continued operation during power mains interruptions, it is recommended that the equipment be powered from an uninterruptible power supply or a battery.
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	30 A/m	30 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.

* U_T is the a.c. mains voltage before application of the test level.

1.9.3 Immunity—RF interference

The following table provides guidance and Stereotaxis’ declaration on **electromagnetic immunity** regarding radiofrequency (RF) for the equipment:

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment—guidance
<p>Conducted RF IEC 61000-4-6</p> <p>Radiated RF IEC 61000-4-3</p>	<p>3 Vrms 150 kHz to 80 MHz</p> <p>3 V/m 80 mHz to 2.7 GHz</p>	<p>3 V</p> <p>3 V/m</p>	<p>Portable and mobile RF communications equipment should be used no closer to any part of the equipment, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.</p> <p>Recommended separation distance</p> <p>$d = 1.2\sqrt{P}$</p> <p>$d = 1.2\sqrt{P}$ 80 MHz to 800 MHz</p> <p>$d = 2.3\sqrt{P}$ 800 MHz to 2.5 GHz</p>
<p>Proximity Fields to RF Wireless Communications</p>	<p>27V/m 380-390Mhz</p> <p>28V/m 430-470Mhz 800-960Mhz 1700-1990Mhz 2400-2570Mhz</p> <p>9V/m 704-787Mhz 5100-5800Mhz</p>	<p>27V/m</p> <p>28V/m</p> <p>9V/m</p>	<p>where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in meters (m).</p> <p>Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey,^a should be less than the compliance level in each frequency range.^b</p> <p>Interference may occur in the vicinity of equipment marked with the following symbol:</p> <div style="text-align: center;">  </div>

Note 1: At 80 MHz and 800 MHz, the higher frequency range applies.

Note 2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects, and people.

- a** Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast, and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the equipment is used exceeds the applicable RF compliance level above, the equipment should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the equipment.
- b** Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.

1.9.4 Separation distances

The equipment is intended for use in the electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the equipment can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the equipment as recommended in the following table, according to the maximum output power of the communications equipment.

The table provides **recommended separation distances** between portable and mobile RF communications equipment and the equipment.

Rated maximum output power of transmitter* W	Separation distance according to frequency of transmitter m		
	150 kHz to 80 MHz $d = 1.2\sqrt{P}$	80 MHz to 800 MHz $d = 1.2\sqrt{P}$	800 MHz to 2.5 GHz $d = 2.3\sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

* For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

Note 1: At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

Note 2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects, and people.



WARNING: Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of the equipment, including cables specified by Stereotaxis, Inc.



Note: If there are Electromagnetic Compatibility (EMC) issues with the equipment, please contact Stereotaxis Service. Otherwise, there are no specific service requirements to maintain EMC integrity.

2 Hardware

The iCONNECT System gathers impedance values in relation to electrode connections. The unaltered values are then sent through the iCONNECT Electronics Hub to *Navigant* where impedance is measured and contact is determined.

2.1 Electronics Hub

The iCONNECT Electronics Hub (Figure 1) has connection points for catheter cables and mapping system/ablation generators. The red port is for the catheter cable and the blue port is for the mapping System or ablation generator cable.

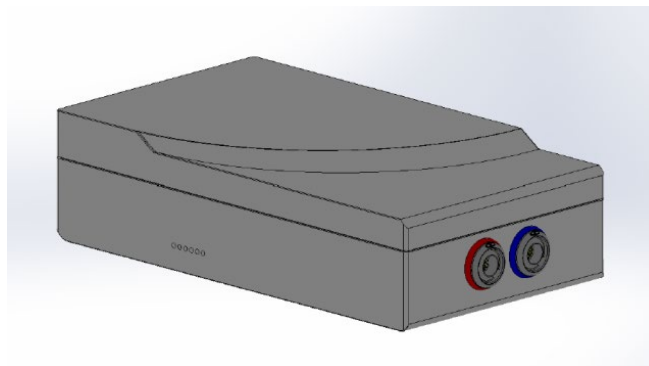


Figure 1. iCONNECT Electronics Hub

2.2 Connection Cables

The iCONNECT System offers use of two sets of connection cables (Figure 2), one for the connection between iCONNECT and the catheter, *Catheter Cables*, and one for connection between iCONNECT and the mapping and/or ablation system, *System Cables*. Catheter cables are provided sterile, with the ability to be resterilized. See **Table 1** and **Table 2** for a list of catheter and system cables and possible connections.

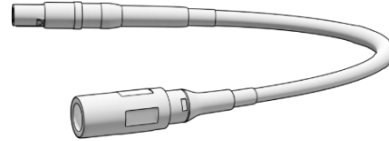


Figure 2. Connection Cable

Table 1. Catheter Cables (Sterile)

Connection	Catalogue Number
iCONNECT – Celsius	001-009071-1
iCONNECT – Magnoflush	001-009073-1

Table 2. System Cables (Non-sterile)

Connection	Catalogue Number
iCONNECT – CARTO	001-009075-1
iCONNECT – Navistar Adapter	001-009081-1
iCONNECT – Maestro	001-009077-1
iCONNECT – Stockert	001-009078-1
iCONNECT – Ampere	001-009079-1
iCONNECT – HAT500	001-009080-1

2.3 Installation

Below are instructions for connecting the iCONNECT Electronics Hub (Figure 3) to the catheter, mapping system, and/or ablation generator. Refer to **Figure 3** and **Figure 4** while following the steps.

Installing the iCONNECT System is a joint venture between the customer and Stereotaxis. **Step 1** must be performed by a Stereotaxis Field Service representative. All subsequent steps can be carried out by the customer.

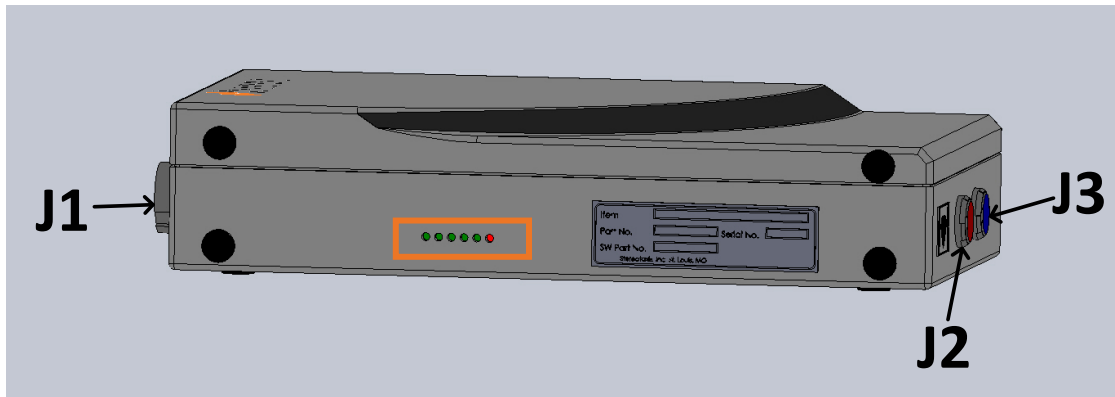


Figure 3. iCONNECT Electronics Hub with connector and LED location identifiers

1. The Stereotaxis Field Service representative will connect the Ethernet cable (P/N 072-011422-1) from A15-J22 of the MNS cabinet to **J1** of the electronics hub. The Ethernet cable provides power and communication for the iCONNECT System.
2. To connect the catheter cable, insert the connector of the catheter cable in the red **J2** socket of the electronics hub and in the ablation catheter in accordance with the information on the cable plugs.
3. If using a Navistar Adapter Cable, insert the connector of the cable in the red **J2** socket of the electronics hub. The opposite end will connect to the Biosense Webster Interface Cable **C5MHNAVMHS**. The free, unconnected end of the Interface Cable connects to the NAVISTAR® RMT Catheter or NAVISTAR® THERMOCOOL® RMT Catheter.
 - a. It is important to note that if a Navistar Adapter Cable is used, an Interface Cable is needed and must be ordered through Biosense Webster.
4. To connect the system cable, insert the connector of the system cable in the blue **J3** socket of the electronics hub and in the RF ablation device or the external connection box for the mapping system.
5. Verify the LEDs are correct according to **Table 3**.



Note: For emphasis, the location of the LEDs is outlined in an orange rectangle in Figure 3.

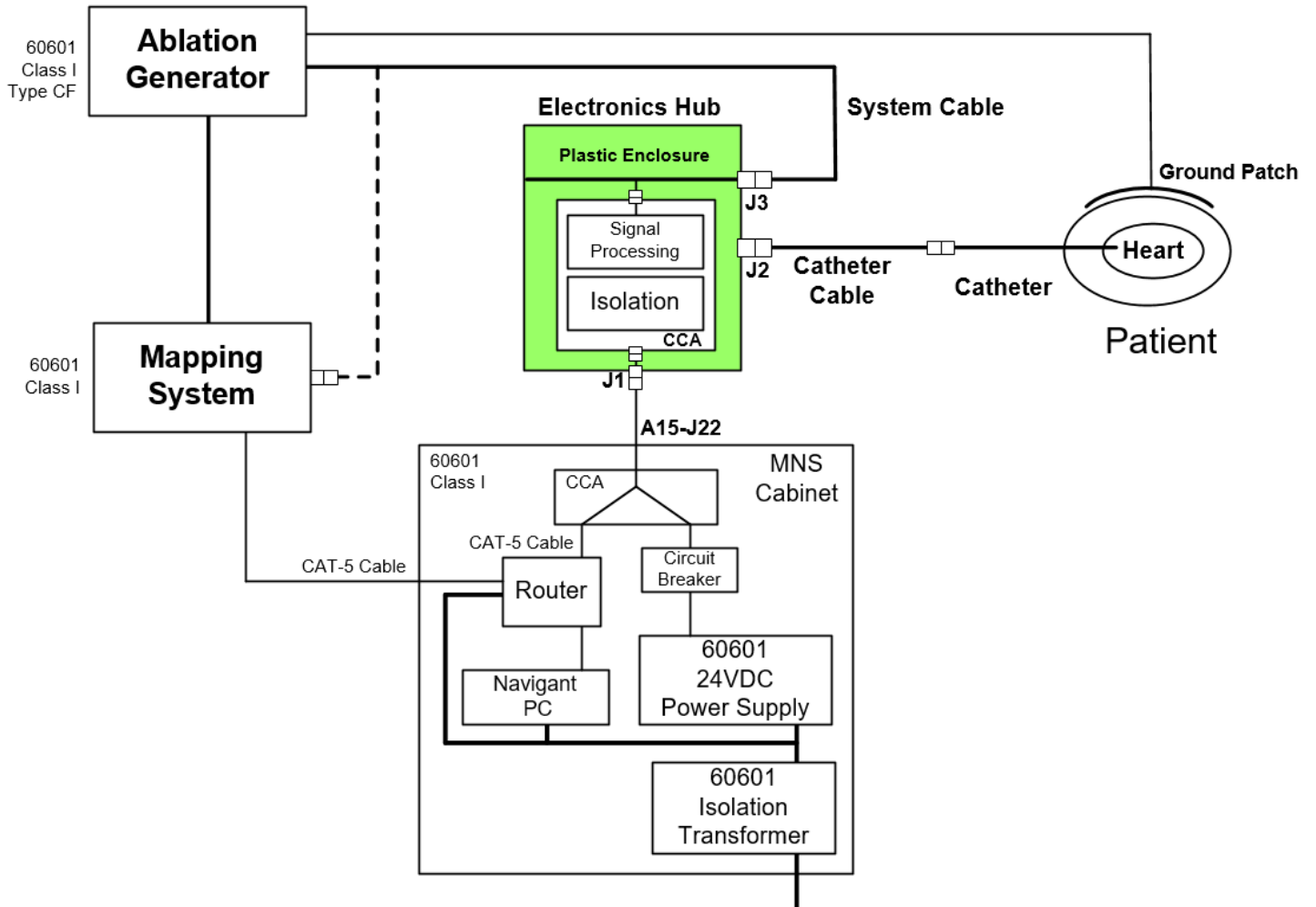


Figure 4. iCONNECT System hardware connectivity

After completing all connections (Figure 4), the external LEDs (Figure 5) on the iCONNECT Electronics Hub box should light as indicated in **Table 3**.



Figure 5. iCONNECT System LEDs (example)

Table 3. iCONNECT System LED indicators

LED	Action
1	Red LED: If lit, indicates error.
2	Starts blinking after a few seconds, indicating processor is active.
3	Final green LED to light, indicating network connectivity to <i>Navigant System</i> .
4	Should light when iCONNECT Ethernet cable is connected.
5	Should light when iCONNECT Ethernet cable is connected.
6	Should light when iCONNECT Ethernet cable is connected.

3 Software

The iCONNECT System is accompanied by software enhancements that measure the level of tip-to-tissue contact and indicate hardware status. iCONNECT communicates with *Navigant* which indicates the level of contact.

3.1 Contact Indication

To indicate contact, the starburst has three states:

- **Optimal Contact** (Figure 6, left)
- **In Contact** (Figure 6, right)
- **No Contact**: No starburst

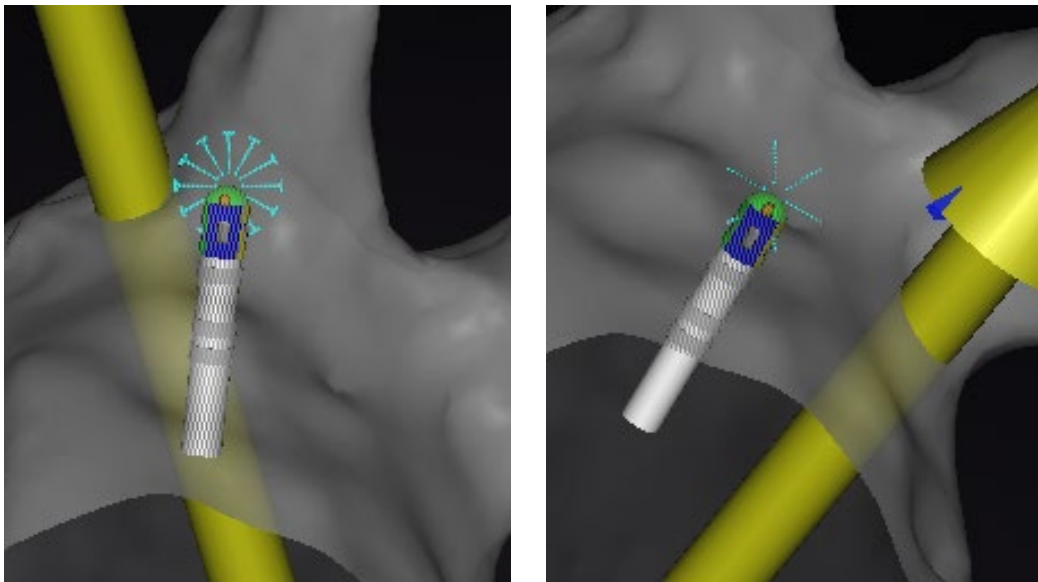


Figure 6. Examples of starburst in **Optimal Contact** (left) and **In Contact** (right)

When the catheter is in contact with the cardiac wall, the *Navigant* screen displays a “starburst” effect, as indicated (②) in **Figure 7**. The *Navigant* screen displays a contact graph in the lower right corner (① in **Figure 7**) to provide a history of cardiac contact.

This graph, or tracing, will be

- *blue* during **Optimal Contact**
- *dashed* during **In Contact**
- *gray* during **No Contact**

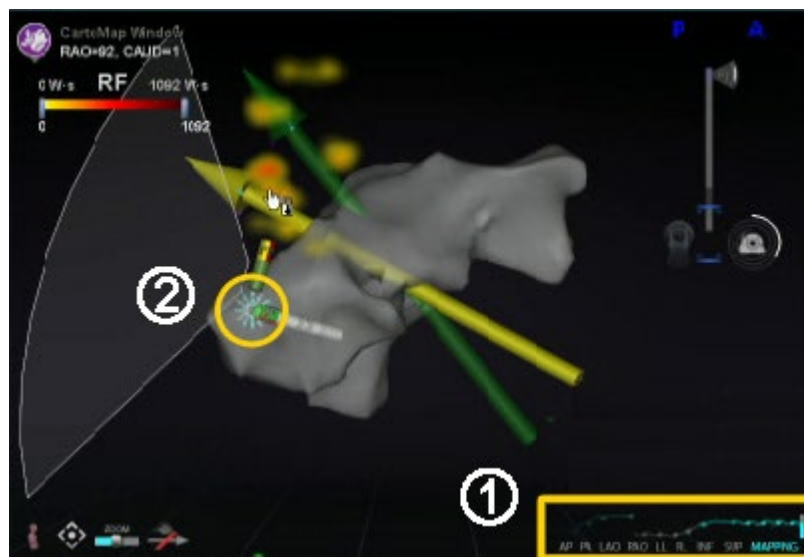


Figure 7. Contact graph on *Navigant* display (gold box outline)



Note: The gold circle and rectangle in Figure 7 are present for emphasis. They do not appear on the *Navigant* screen.

3.2 Impedance Graph

When the Impedance Graph feature is enabled, an iCONNECT System is connected, and a compatible catheter is positioned in the heart, a graph is shown at the bottom of *Navigant's* 3D window.

The Impedance Graph overlay displays two tracings (emphasized by a red rectangle in Figure 8). The first purple tracing is the resistance between the tip and electrode 3. The green tracing is the resistance between electrode 2 and 4.

The total duration of the impedance graph is 120 seconds. The right side of the graph reflects impedance in real time, at the tip of the catheter. The dashed line at the bottom of the impedance graph represents the baseline (the range of the y-axis, measured in ohms, being 55 above baseline to 5 below baseline). The baseline is determined in real time and changes according to incoming data. The scale of the graph and y-axis are always relative to the current baseline.

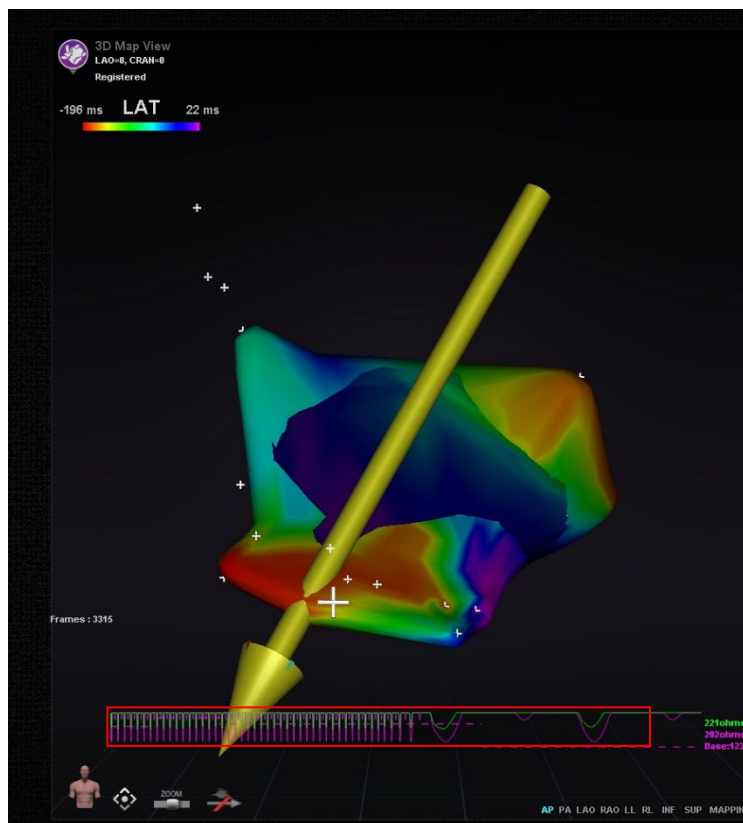


Figure 8. Impedance Tracings

To enable the Impedance Tracing overlay, click on the *Navigant* start button and open the Settings dialog by choosing the settings menu item (Figure 9). Select the 3D window and turn on the iCONNECT Impedance Graph option. (Without connection to iCONNECT, the impedance graph will not display.)

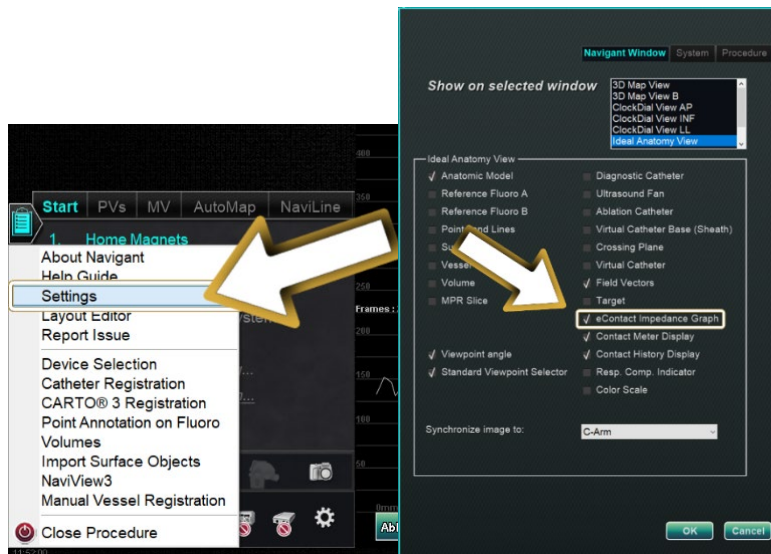


Figure 9. Settings to enable Impedance graphs

3.3 Hardware Status Panel Display

The *Navigant* hardware status indicator bar shows an additional icon when the iCONNECT System feature is present (Figure 10). As shown in **Figure 11**, the iCONNECT System icon has three possible statuses.

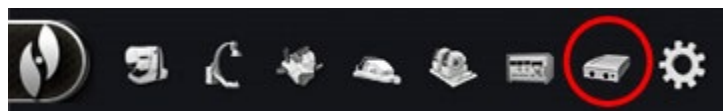


Figure 10. *Navigant* hardware status indicator bar with iCONNECT System icon



Figure 11. iCONNECT System status icon in three states

Table 4 lists examples of the **Warning** and **Error** iCONNECT System states and the possible reasons or reported reasons:

Table 4. iCONNECT System states

State	Reason
Warning	Contact data is out of range (alert is cleared when contact data returns to correct range)
Error	<i>Navigant</i> System not connected to iCONNECT (alert is revoked when <i>Navigant</i> System connects to iCONNECT)
Error	iCONNECT reporting error (alert is revoked when device reports that error has been cleared)

4 Maintenance

4.1 Cleaning

Do not autoclave or submerge the iCONNECT Electronics Hub. The iCONNECT Electronics Hub and cables are designed to be wiped down with a cloth or swab dampened in pH neutral, EPA-approved hospital grade solutions.

4.2 Sterilization of Cables

The iCONNECT Catheter Cables will be supplied sterile. Please refer to the respective catheter cable IFU for more information regarding sterility and resterilization methods of the cables.

5 Troubleshooting

If you need assistance with any of the solutions in the following table, call the TeleRobotic Support Team (TST) at one of the numbers in the following section.

Problem	Solution	
iCONNECT System unit LED indicator errors (see Figure 5 for LED locations)	> 0 and < 3 leftmost LEDs are lit.	Call the TST for Field Service evaluation.
	All 3 leftmost LEDs are on and LED 3 is <i>not</i> blinking.	Call the TST for Field Service evaluation.
	All 3 leftmost LEDs are on and LED 3 is blinking, but LED 4 is not lit.	Further investigation is required. Call the TST.
	Red LED is lit.	An error has occurred. Unplug the Ethernet cable and plug it back in. If the problem persists, call the TST.
iCONNECT warning and errors (in relation to the Navigant System)	iCONNECT unit reports contact data that is out of range. <i>Navigant System</i> indicates a persistent warning alert by way of the iCONNECT System status icon.	Check the catheter connections. If the warning alert fails to clear, call the TST.
	iCONNECT unit reports board error. <i>Navigant System</i> indicates a persistent error alert by way of the iCONNECT status icon.	The alert is revoked when the unit reports the error has been cleared. If the unit fails to revoke the error, call the TST.
	<i>Navigant System</i> is not connected to iCONNECT. <i>Navigant System</i> indicates a persistent error alert by way of the iCONNECT status icon.	Connect the <i>Navigant System</i> to the <i>e-Contact Module</i> unit. The alert should clear, but if the alert fails to clear, call the TST.



Note: For questions regarding non-Stereotaxis products, refer to the respective User Guides or IFUs.



If iCONNECT is still not operating properly after troubleshooting and ensuring all connectors are attached correctly, call the TeleRobotic Support Team for further help:

www.stereotaxis.com

4320 Forest Park Avenue, Suite 100

St. Louis, MO 63108

1-314-678-6200

1-866-269-5268 (Toll free in US)

Appendix A: CARTO System Cable and Navistar Adapter Cable

Instructions for Use