

# Nurse Guidelines

## Fluorescence Guided Imaging Preparation & Troubleshooting

### Fluorescence Guided Imaging & Indocyanine Green

#### General Concepts

- Fluorescence guided imaging (FGI) is a surgical tool implemented to improve the operating field imagery feedback and provided real-time guidance of target anatomy.
- This technique operates through the fluorescent dye, indocyanine green (ICG).
- Indocyanine green (ICG) is a dye and optical imaging agent indicated for fluorescence guided surgery, which includes but is not limited to: Ophthalmic-, foregut-, bariatric-, hepatobiliopancreatic-, colorectal-, endocrine-, plastic-, gynecologic-, cardiothoracic-, pediatric-, genitourinary-, oncologic-, and neurosurgery.
- Its applications focus on the optical enhancement of micro- and macro-vasculature, blood flow, target tissue perfusion, and tumoral and sentinel lymph node isolation and identification in the presence of malignancies.
- The technique can be performed via hand-held devices for open procedures, or totally endoscopic devices for laparoscopic or robotic-assisted procedures.

#### Pharmacokinetics & Posology

- The ICG dye is relatively innocuous, and its tandem-device applications render no radiation. Nevertheless, the operating room nurse and other personnel should be aware of its general pharmacology.
- The standard presentation for the United States and the America's is a powdered vial of 25 mg of ICG. See the '*Precautions*' section.
- The metabolism of ICG is hepatic, with a percentage disappearance rate in healthy subjects of 18-24% per minute.
- Normal biological half-time is 2.5-3.0 minutes.
- **Adults max dose**
  - Do not exceed a maximum total dose of 2 mg/kg
- **Infants, children, and adolescents max dose**
  - Do not exceed a maximum total dose of 2 mg/kg
- **Contraindications**
  - Relative contraindication: ICG contains sodium iodide and should be used with caution in patients who have a history of allergy to iodides.

- **Warnings**

- Anaphylactic deaths have been reported following ICG administration
- Overall incidence of 0.05%
  - Reactions: Anaphylactic or urticarial reactions have been reported in patients with or without a history of allergy to iodides. If such reactions occur, treatment with the appropriate agents (e.g., epinephrine, antihistamines, and corticosteroids) should be administered.

- **Precautions**

- ICG is unstable in aqueous solution and must be used within 6 hours.
- The dye should be diluted and flushed, when indicated, with sterile water to avoid precipitation of the dye within the plasma.
- The dye is stable in plasma and whole blood so that samples obtained in discontinuous sampling techniques may be read hours later.
- Sterile techniques should be used in handling the dye solution as well as in the performance of the dilution curves.
- The un-reconstituted (powder) dye may cling to the vial or lump together because it is freeze-dried in the vials.

- **Drug interactions**

- Heparin preparations containing sodium bisulfate reduce the absorption peak of ICG in blood and, therefore, should not be used as an anticoagulant for the collection of samples for analysis.

- **Carcinogenesis, Mutagenesis, Impairment of Fertility**

- No studies have been performed to evaluate the carcinogenicity, mutagenicity, or impairment of fertility.

- **Pregnancy**

- Teratogenic Effects: Pregnancy Category C - can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity.

- **Lactation**

- It is not known whether this drug is excreted in human milk.
- Caution should be exercised when ICG is administered to a nursing individual.

## Preoperative Precautions

### Circulator Nurse Tasks

#### ***Complete prior to patient entry into the operating room***

1.	<i>OR table position:</i> Ensure position required for the procedure being performed. <b><i>IMPORTANT:</i></b> Account for external arms or table extensions for device placement (e.g. laparoscope or hand-held device stand).
2.	<i>Power sources:</i> Check that all power sources are connected, and device units are switched 'ON'.
3.	<i>Video connectors:</i> Check that screen displays and device being used are connected and transmitting signal correctly. <b><i>IMPORTANT:</i></b> Account for required cables (e.g. HDMI-to-HDMI, DVI-to-DVI, VGA-to-VGA, etc.).
4.	Ensure that video/imaging documentation sources are operational. Capture a small recording and an image to confirm.
5.	Complete ICG time-out checklist.

#### ***Scrub Tech/RN Sterile Equipment Tasks***

1.	Check for drapes, external arms or stands associated with the intended device used for the procedure.
2.	Check for sterile cup, syringes, and needles availability in case of tandem ' <i>in-field</i> ' administration of ICG.
3.	Check laparoscope or hand-held device for clarity and vision.
4.	In the case of further ' <i>in-field</i> ' dilutions of the dye, check for availability of sterile water.



## Intraoperative Troubleshooting

### **NO PICTURE ON MONITOR**

Camera control or other components (displays, media hub, etc.) not on.

Make sure all power sources are activated and correctly plugged in.

### **NO PICTURE ON MONITOR**

Cable connectors between device control unit and/or monitor not attached properly.

Cable should run from output on device to input on primary monitor. Use compatible cables for device and light source.

### **NO PICTURE ON MONITOR**

Input selection on monitor or video peripherals not (displays, media hub, etc) selected.

Adjust input selection.

# Operating Room Checklist



Date		Medical Record Number	
Lastname, Name			
D.O.B.		Weight (kg)	
Procedure			
ICG contraindications	Iodine allergies <input type="checkbox"/> Mercury allergies <input type="checkbox"/> Pregnancy <input type="checkbox"/> Lactation <input type="checkbox"/>		

## ICG TIME-OUT

	Yes	No
Confirmed patient identity	<input type="checkbox"/>	<input type="checkbox"/>
Is this a pediatric case	<input type="checkbox"/>	<input type="checkbox"/>
Confirmed procedure	<input type="checkbox"/>	<input type="checkbox"/>
NIR device in room	<input type="checkbox"/>	<input type="checkbox"/>
Device ready for recording including images and video	<input type="checkbox"/>	<input type="checkbox"/>
Sterile covers available for intended device	<input type="checkbox"/>	<input type="checkbox"/>
ICG dye in room	<input type="checkbox"/>	<input type="checkbox"/>
Confirmed ICG dosage	<input type="checkbox"/>	<input type="checkbox"/>
Further dilution required	<input type="checkbox"/>	<input type="checkbox"/>
Sterile water readily available for reconstitution AND flushing	<input type="checkbox"/>	<input type="checkbox"/>
Confirm site of administration (e.g.: Peripheral, intradermal, etc)	<input type="checkbox"/>	<input type="checkbox"/>
Industry contacts available in case of device/dye troubleshoot	<input type="checkbox"/>	<input type="checkbox"/>
Verbal confirmation for administration prior to local analgesia administration	<input type="checkbox"/>	<input type="checkbox"/>
Anaesthesia confirmed	<input type="checkbox"/>	<input type="checkbox"/>
Special considerations/Additional comments		