HEPZATO KIT contains
HEPZATO (melphalan) for Injection/
Hepatic Delivery System (HDS)

Percutaneous Hepatic Perfusion Procedure

REMS Didactic Modules Training Program Modules 1 – 9

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Deleath

HEPZAT® KIT™





HEPZATO KIT REMS

Didactic Modules Training Overview

Delcath





Delcath Systems, Inc.'s detailed description of the clinical use of the **HEPZATO KIT** can be found in the Instructions For Use manual (IFU) and in the US Prescribing Information.

The IFU and US Prescribing Information should be thoroughly reviewed prior to training.



HEPZATO KIT REMS Didactic Modules Training Overview





This Didactic training is intended to educate the Healthcare Setting staff on the Risk Evaluation and Mitigation Strategy (REMS) and the safe and effective use of HEPZATO (melphalan) and the HEPZATO KIT Hepatic Delivery System (HDS).



HEPZATO KIT is indicated as a liver-directed treatment for adult patients with uveal melanoma with unresectable hepatic metastases affecting less than 50% of the liver and no extrahepatic disease, or extrahepatic disease limited to the bone, lymph nodes, subcutaneous tissues, or lung that is amenable to resection or radiation.



HEPZATO KIT should only be utilized for percutaneous hepatic procedures (PHP) performed by qualified personnel trained in accordance with the IFU and the PI.

HEPZATO KIT REMS Didactic Modules Training Overview



This training is comprised of nine Didactic Modules



Training requirements are established for each specialist role of the PHP Procedure team



All PHP Procedure team members are required to complete the didactic module that provides an overview for their role in the procedure

Module #	Module	IR	Anesthesiologist	Perfusionist
1	REMS Program Details and Requirements	Required	Required	Required
2	HEPZATO Overview	Required	Required	Required
3	Treatment Planning	Required	Required	Required
4	Perfusion Circuit	Optional	Required	Required
5	Drug Preparation & Injection	Optional	Optional	Required
6	Procedure Day	Required	Required	Required
7	Catheterization & Drug Infusion	Required	Optional	Required
8	Hemodynamics Management	Optional	Required	Optional
9	Post Procedure	Optional	Required	Optional

The Oncologist and other staff members may also complete Didactic Module training but are not required to complete this training.

HEPZATO KIT Clinical Treatment Team

HEPZATO KIT Clinical Treatment Team

PHP PROCEDURE TEAM (REQUIRES REMS TRAINING)

- Interventional Radiologist (IR)
 IR or Surgical Oncologist must be the LEADER during the procedure and communication is paramount
- Anesthesiologist (AN)
- Perfusionist (PF)

OTHER HEPZATO KIT CLINICAL TREATMENT TEAM MEMBERS (DO NOT REQUIRE REMS TRAINING)

- Oncologist (Medical and/or Surgical; SO/MO)
 - Must commit to managing the patient before and after the procedure
 - May be involved during the procedure
- Pharmacist
- Registered Nurse (RN)
- Interventional Radiology Staff
- Intensivist or Critical Care Specialist

 Manages patient immediately post-procedure along with the SO/MO

Each health care professional must communicate and collaborate to ensure that each of the individual procedures are appropriately timed.







MODULE 1

Risk Evaluation & Mitigation Strategy (REMS) Program Details and Requirements

HEPZATO KIT REMS Didactic Modules Training

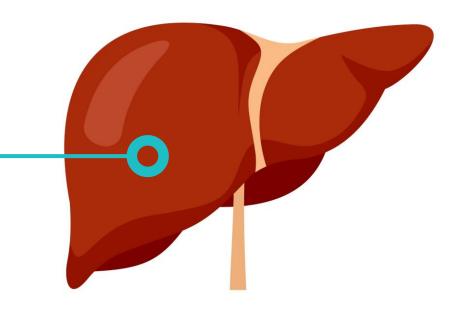


This educational didactic module contains information on serious adverse events associated with the use of HEPZATO KIT, including the risks of severe peri-procedural complications including hemorrhage, hepatocellular injury, and thromboembolic events. This is not a comprehensive list of adverse events associated with HEPZATO KIT.

Please refer to the prescribing information for a comprehensive list of adverse events.

HEPZATO Indication

HEPZATO is indicated as a liver-directed treatment for adult patients with uveal melanoma with unresectable hepatic metastases affecting less than 50% of the liver and no extrahepatic disease, or extrahepatic disease limited to the bone, lymph nodes, subcutaneous tissues, or lung that is amenable to resection or radiation.



What is a REMS?



A Risk Evaluation and Mitigation Strategy (REMS) is a program required by the FDA to manage known or potential serious risks associated with a drug product. The FDA has determined that a REMS is necessary to ensure that the benefits of HEPZATO KIT outweigh its risks.



The goal of the HEPZATO KIT REMS is to mitigate the risks of severe peri-procedural complications including hemorrhage, hepatocellular injury, and thromboembolic events associated with HEPZATO KIT.

HEPZATO KIT REMS Summary



HEPZATO KIT is only available through the HEPZATO KIT REMS



Healthcare settings
that dispense
HEPZATO KIT must
be certified in the
REMS



PHP procedure team members that perform procedures with HEPZATO KIT must be trained

PHP Procedure Team







HEPZATO KIT Clinical Treatment Team



HEPZATO KIT Clinical Treatment Team - Others









Percutaneous Hepatic Perfusion (PHP) Procedure Team



The percutaneous hepatic perfusion procedure team(s) must include healthcare providers with expertise in interventional radiology, anesthesiology, and perfusion in accordance with the Instructions for Use.

Healthcare Setting Certification Requirements

As a condition of certification, the Healthcare Setting must:



Have a percutaneous hepatic perfusion procedure team(s) that must include healthcare providers with expertise in interventional radiology, anesthesiology, and perfusion as described in the Instructions for Use.



Have the following on-site: interventional radiology suite or operating room with fluoroscopy and with resuscitation personnel, equipment, and medications



Authorized Representative Requirements



The Healthcare Setting must designate an **Authorized Representative (AR)** to carry out the certification process and oversee implementation and compliance with the REMS requirements on behalf of the healthcare setting.

To become certified, the Authorized Representative must:



Carry out the certification process and oversee implementation and compliance with the REMS requirements on behalf of the healthcare setting.



Review HEPZATO KIT REMS Prescribing Information, Instructions for Use, **Program Overview**, and this **Didactic Modules**



Enroll in the REMS Program by completing and submitting the **Healthcare Setting Enrollment Form** to the REMS



Establish processes and procedures to ensure new members of the PHP procedure team are trained and successfully complete the Proctorship training and complete and submit the **Criteria for Procedural Competency Checklist** to the REMS.

Who Can Be an Authorized Representative?

An Authorized Representative at the Healthcare Setting can be a:



Percutaneous Hepatic Perfusion (PHP) Team Requirements

The PHP Procedural Team will be required to:



Review the HEPZATO
Prescribing Information
and HEPZATO KIT
Instructions for Use



Review the Program
Overview, this Didactic
Module and undergo the
Preceptorship Training
provided by Delcath
Systems, Inc.



Successfully complete the Proctorship and Criteria for Procedural Competency Checklist

Before HEPZATO KIT Administration

Before administering HEPZATO KIT, the Healthcare Setting must:



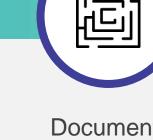
Obtain an authorization to dispense by contacting the REMS Coordinating Center to verify the PHP Procedure Team is qualified using the **Procedure**Team Qualification Form



During HEPZATO KIT Administration

During and for a minimum of 72 hours after the PHP procedure, the Healthcare Setting must:







Assess the patient for severe peri-procedural complications associated with HEPZATO KIT.

Document severe peri-procedural complications using the **Severe Peri-Procedure-related Complications Adverse Events Documentation Form** and submit to the REMS.

Follow up may be virtual or via telephone in patients discharged prior to 72 hours

Maintain Healthcare Setting Certification

To maintain the ability to dispense the HEPZATO KIT, a Healthcare Setting must:



Have a PHP procedure team(s) that must include healthcare providers with expertise in interventional radiology, anesthesiology, and perfusion



Have PHP team members who have each performed one PHP procedure in the first six months following completion of training, a second procedure in the next six months, and at least two procedures annually thereafter



Have the following on-site: interventional radiology suite or operating room with fluoroscopy and with resuscitation personnel, equipment, and medications



Have a new Authorized Representative enroll in the REMS by completing the **Healthcare Setting Enrollment Form** and submitting it to the REMS

Healthcare Setting Responsibilities

At all times, the Healthcare Setting must:



Maintain records of each PHP team members training



Maintain records of the PHP procedures performed with HEPZATO KIT and the associated PHP procedure team members' participation



Maintain records that indicate the processes and procedures are in place and being followed



Comply with audits carried out by Delcath Systems, Inc., or a third party acting on behalf of Delcath Systems, Inc., to ensure that all REMS specific processes and procedures are in place and being followed

HEPZATO KIT REMS Materials Summary

Program Overview	 Reviewed by the Authorized Representative and the PHP procedure team members Educates on the key risks, management of the risks, and REMS requirements 		
Didactic Modules	 Reviewed by the Authorized Representative and the PHP procedure team members Educates on the key risks, management of the risks, and REMS requirements 		
Healthcare Setting Enrollment Form	Completed and submitted by a designated Authorized Representative Agreement to comply with the REMS requirements		
Criteria for Procedural Competency Checklist	Completed by the PHP procedure team members as part of the Proctorship Retained on-site as documentation of training and submitted to the REMS		
Procedure Team Qualification Status Form	 Completed and submitted by the Authorized Representative and/or Healthcare Setting staff to obtain authorization to dispense each HEPZATO KIT to verify the PHP procedure team is qualified 		
Severe Peri-Procedure- Related Complications Adverse Events Documentation Form	 Completed and submitted by the Healthcare Setting to report any severe peri-procedural complications associated with HEPZATO KIT during and after administration for at least 72 hours 		
REMS Program Website	Dedicated resource for REMS information		

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HEPZATO KIT Procedural Overview







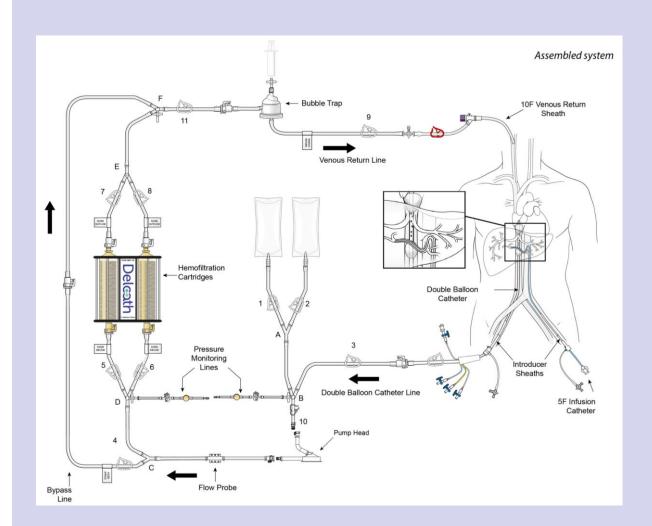




HEPZATO KIT is a closed circuit of catheters and filters utilized to deliver a chemotherapeutic agent (melphalan) directly with simultaneous filtration of hepatic venous blood during drug infusion and washout, which results in locoregional delivery of a relatively high melphalan dose but also reduced systemic exposure.

There are risks including peri-procedural complications that may be severe and or life threatening

The prominent risks include hemorrhage, hepatocellular injury, and thromboembolic events



Severe Peri-Procedural Complications

Hemorrhage, hepatocellular injury, and thromboembolic events have been observed when HEPZATO has been administered via hepatic intra-arterial administration.



Administration of HEPZATO requires general anesthesia and extracorporeal bypass of circulation which may cause life threatening or fatal adverse effects



Ensure the patient is euvolemic but do not overhydrate the patient.



Monitor for these periprocedural complications during the procedure and for at least 72 hours following the procedure.

Contraindications

HEPZATO KIT is contraindicated in patients with:

Active intracranial metastases or brain lesions with a propensity to bleed

Liver failure, portal hypertension, or known varices at risk for bleeding

Surgery or medical treatment of the liver in the previous 4 weeks

Active cardiac conditions including, but not limited to, unstable coronary syndromes (unstable or severe angina or myocardial infarction), worsening or new-onset congestive heart failure, significant arrhythmias, or severe valvular disease

History of allergies or known hypersensitivity to melphalan or a component or material utilized within the HEPZATO KIT including natural rubber latex, heparin, and severe hypersensitivity to iodinated contrast not controlled by antihistamines and steroids

Warnings and Precautions (1)

Hemorrhage, hepatocellular injury, and thromboembolic events have been observed when melphalan has been administered via hepatic intra-arterial administration of HEPZATO. Administration of HEPZATO requires general anesthesia and extracorporeal bypass of circulation which may cause life threatening or fatal adverse effects. Ensure the patient is euvolemic but do not overhydrate the patient. Monitor for these peri-procedural complications during the procedure and for at least 72 hours following the procedure.

To mitigate the risk of thromboembolic events, administer anticoagulation as described in the IFU during the procedure.

Warnings and Precautions (2)

Due to the risk of bleeding, do not use in patients with uncorrectable coagulopathies and delay treatment with the HEPZATO KIT for at least 4 weeks after surgery or other medical procedure involving the liver. Platelets and clotting factors may be removed during the HEPZATO KIT procedure. Monitor platelets and coagulation parameters as described in the IFU. If life-threatening bleeding occurs during the procedure, reverse anticoagulation as described in the IFU and correct coagulopathy as appropriate. Discontinue anticoagulation with warfarin or other oral anticoagulants prior to the procedure until hemostasis has been restored after the procedure and no bleeding complications have been observed. Refer to the Prescribing Information of the anticoagulant agent for bridging recommendations for anti-coagulation prior to surgical procedures. Discontinue drugs affecting platelet function such as aspirin, non-steroidal anti-inflammatory drugs, or other anti-platelet drugs one week before the procedure.

Warnings and Precautions (3)

Patients with abnormal hepatic vascular (especially arterial supply) or biliary (especially reimplantation of bile duct) anatomy or gastric acid hypersecretion syndromes may be at increased risk of peri-procedural complications or other severe adverse reactions. Screen patients for a history of prior surgeries involving the bile duct to assess whether the patient is an appropriate candidate for HEPZATO KIT and monitor patients for adverse reactions following HEPZATO KIT administration.

Procedure-related reductions in blood pressure including severe hypotension can occur during the HEPZATO KIT procedure. Closely monitor blood pressure during the procedure. Patients may require fluid support and vasopressors. To reduce the risk of severe hypotension, temporarily discontinue ACE-inhibitors, calcium channel blockers, or alpha-1-adrenergic blockers for at least 5 half-lives prior to treatment with the HEPZATO KIT. If necessary, use other short-acting antihypertensive drugs to manage blood pressure during the peri-procedure period.

Warnings and Precautions (4)

Hematologic adverse reactions, including thrombocytopenia, anemia, and neutropenia have been reported in patients treated with HEPZATO. The risk of hematologic adverse reactions may be increased in patients who have received prior chemotherapy, bone irradiation, or who have compromised bone marrow function.

Monitor patients for severe infections, bleeding, and symptomatic anemia. Only administer HEPZATO in patients with platelets >100,000/microliter, hemoglobin ≥10.0 gm/dL and neutrophils >2,000/microliter. Administer transfusions or growth factors as appropriate

Warnings and Precautions (5)

Hypersensitivity reactions, including anaphylaxis, have occurred in approximately 2% of patients who received an intravenous (IV) formulation of melphalan. These reactions with melphalan are characterized by urticaria, pruritus, edema, skin rashes, and in some patients, tachycardia, bronchospasm, dyspnea, and hypotension. Hypersensitivity can occur in patients with or without prior exposure to IV or oral melphalan.

When a hypersensitivity reaction is observed, immediately terminate the hepatic arterial HEPZATO infusion and administer necessary supportive care.

Patients with a history of allergic reactions to iodinated contrast may experience hypersensitivity reactions, including anaphylaxis, during treatment with the HEPZATO KIT. Premedicate patients with a history of allergic reaction to iodinated contrast prior to treatment with HEPZATO KIT. Do not administer HEPZATO KIT in patients with a history of severe allergic reactions or anaphylaxis to iodinated contrast.

Warnings and Precautions (6)

Gastrointestinal adverse reactions including nausea and vomiting, abdominal pain, and diarrhea are common, and occurred in 84% of patients treated with HEPZATO in the FOCUS trial. Administer a proton-pump inhibitor the day prior to and the morning of the procedure. If antiemetic treatment is required, pre-medicate with anti-emetic therapy in subsequent cycles.

Melphalan has been shown to cause chromatid or chromosome damage in humans. Secondary malignancies, including acute nonlymphocytic leukemia, myeloproliferative syndrome, and carcinoma, have been reported in patients with cancer treated with intravenous alkylating drugs including melphalan. Some patients also received other chemotherapeutic agents or radiation therapy. Precise quantification of the risk of acute leukemia, myeloproliferative syndrome, or carcinoma is not possible. Published reports of leukemia in patients who have received oral or IV melphalan (and other alkylating drugs) suggest that the risk of leukemogenesis increases with chronicity of treatment and with cumulative dose.

Warnings and Precautions (7)

Based on animal studies and its mechanism of action, melphalan can cause fetal harm when administered to a pregnant woman. Melphalan is genotoxic, targets actively dividing cells, and was embryolethal and teratogenic in rats. Advise pregnant women of the potential risk to a fetus. Advise females of reproductive potential to use effective contraception during treatment with HEPZATO and for 6 months after the last dose. Advise males with female partners of reproductive potential to use effective contraception during treatment with HEPZATO and for 3 months after the last dose.

Melphalan-based chemotherapy regimens have been reported to cause suppression of ovarian function in premenopausal women, resulting in persistent amenorrhea in approximately 9% of patients. Reversible or irreversible testicular suppression has also been reported.



Adverse Events and Complications

In the multicenter, open label pivotal study, [FOCUS (NCT02678572)], serious adverse reactions occurred in 45% of patients who received HEPZATO.

Serious adverse reactions occurring in ≥ 2% of patients were thrombocytopenia (10%), neutropenia (8%), febrile neutropenia (7%), platelet count decreased (6%), leukopenia (4.2%), cardiac arrest (3.2%), neutrophil count decreased (2.1%), hypoxia (2.1%), pleural effusion (2.1%), pulmonary edema (2.1%), and deep vein thrombosis (2.1%).

Fatal adverse reactions occurred in 3 (3.2%) patients who were treated with HEPZATO; these included cardiac arrest, acute hepatic failure and bacterial peritonitis.

Adverse Event Reporting



Reporting of serious adverse events after administration of HEPZATO KIT is vital for the continued monitoring of the risk/benefit balance of the use of HEPZATO KIT



Healthcare providers must report severe peri-procedural complications including adverse events of hemorrhage, hepatocellular injury, and thromboembolic events using the Severe Peri-Procedure-related Complications Adverse Events Documentation Form to Delcath Systems, Inc. via e-mail at coordinator@HEPZATOKITREMS.com



Healthcare providers are encouraged to report other adverse reactions to Delcath at 1-833-632-0458 or to the FDA at 1-800-FDA-1088 or www.fda.gov/medwatch

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HEPZATO KIT REMS Information

For further information, please visit www.HEPZATOKITREMS.com or call 1-833-632-0457









MODULE 2 HEPZATO KIT Overview



MODULE 2

Purpose and Objectives

The introductory module provides a general overview of **HEPZATO KIT** and PHP procedure including target patient population and indication, contraindications, general safety and precautions, and clinical efficacy/safety information. Training will prepare HCPs for individual education modules based on specialized role.



Understand the mechanism of action of melphalan and the HDS and how these components work as a system.



Understand melphalan clinical pharmacology, the pharmacokinetic profile, and how the system limits melphalan systemic exposure.



Review target patient population and indications, contraindications, precautions, and special populations.



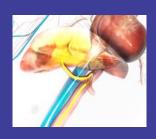
Understand the safety and adverse events profile for HEPZATO KIT and methods to mitigate adverse events.

HEPZATO KIT is a drug/device combination product and is indicated as a liver-directed treatment for adult patients with uveal melanoma with unresectable hepatic metastases affecting less than 50% of the liver with no extrahepatic disease, or extrahepatic disease limited to the bone, lymph nodes, subcutaneous tissues, or lung that is amenable to resection or radiation.

The PHP
Procedure with
HEPZATO KIT
is based on
3 principles



Isolation of the liver circulation through the occlusion of the inferior vena cava caudal and cranial of the hepatic veins.



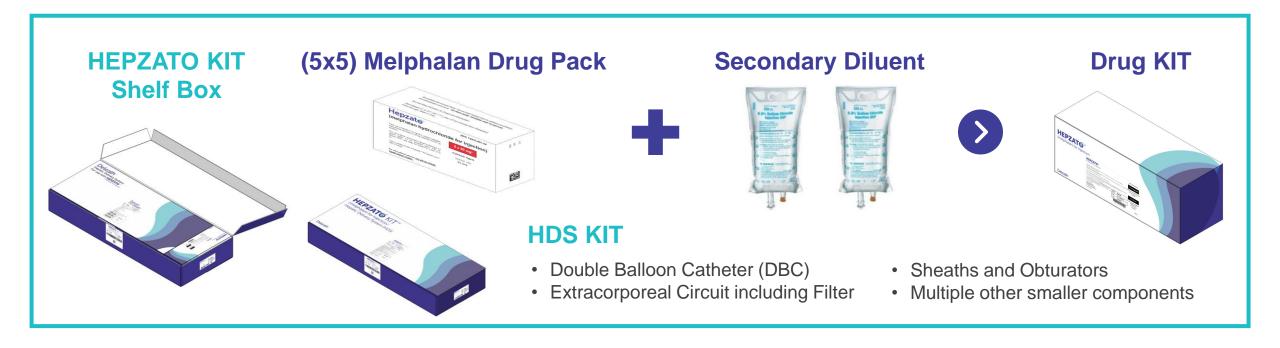
Infusion of high dose melphalan directly into the liver to **saturate** hepatic parenchyma and residing cancer cells.



Filtration of the hepatic circulation to limit melphalan concentration before returning the blood to the systemic circulation thereby reducing melphalan systemic exposure.

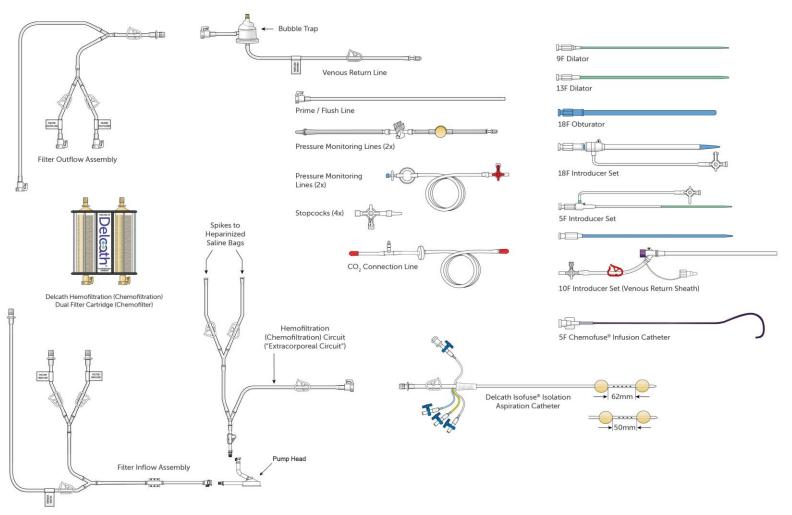
HEPZATO KIT System Components

HEPZATO KIT consists of a closed circuit of catheters and drug specific filters utilized to deliver a chemotherapeutic agent (melphalan) to the hepatic artery and to lower the concentration of the agent in the blood before it is returned to systemic circulation.



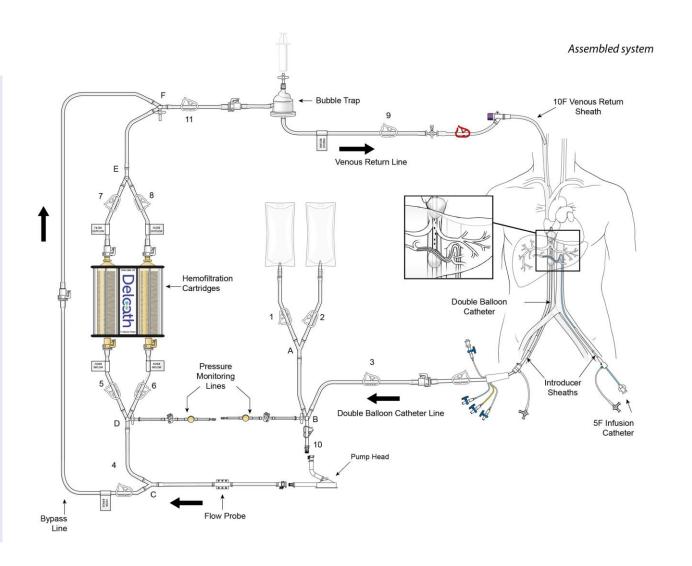
Supplied Disposable Components





Assembled System

Deleath HEPZAT® KIT™ (melphalan) for Injection/ Hepatic Delivery System (HDS) See IFU for **Instructions For Use** full system description



Additional Items needed (not included)

- Bubble Trap holder
- Medtronic Bio-Console 560 Speed Controller System ("Pump")
- Medtronic 560A ("Motor Drive")
- Medtronic Bio-Probe TX50P ("Flow Transducer")
- CO₂ Supply for Priming Dual Filter
- Drug Delivery Disposables:
 - One (1) Medrad 150mL Syringe (Polypropylene (PP)-Barrel & Polyisoprene-Plunger) or equivalent
 - Two (2) Intravenous Administration Set with spike & drip chamber (Polyvinylchloride (PVC)-tubing, Acrylonitrile butadiene styrene (ABS) & Polyethylene (PE)-Drip Chamber & Polycarbonate (PC)-Luer) or equivalent
 - One (1) 48" injector lines (PVC-Tubing & PC-Luer) or equivalent
 - Five (5) 3-way stopcocks (PC-body, High Density Polyethylene (HDPE) or Acetal-Handles) or equivalent
 - Three (3) 20 mL syringes (PP-Barrel & Polyisoprene-Plunger) or equivalent
- Microcatheters (Maximal Distal End OD = 2.8F) for Selective Drug Infusion (at Interventional Radiologist discretion). Select one from Delcath qualified microcatheters listed below:
 - Merit Maestro (Merit Medical Systems, Inc., So. Jordan, UT, USA)
 - BSC Renegade Hi-Flo (Boston-Scientific Corp.; Natick, MA, USA)
 - Terumo Progreat (Terumo Medical Corp., Somerset, NJ, USA)

Healthcare Setting Requirements



The procedure must be performed in an appropriately equipped interventional radiology suite with fluoroscopy or an operating room designed and equipped similarly.

Resuscitation personnel, equipment, and medication must be immediately available.



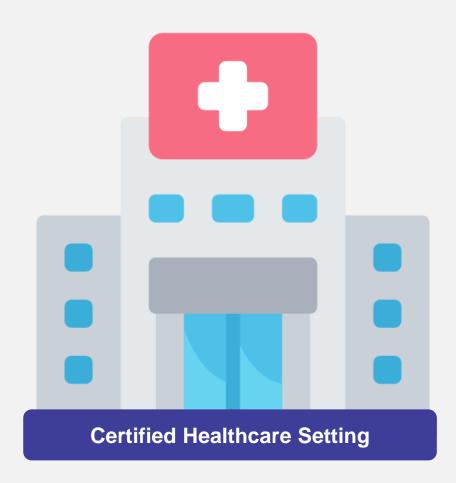
PHP Procedure Team







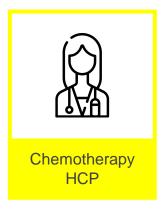
HEPZATO KIT Clinical Treatment Team



HEPZATO KIT Clinical Treatment Team - Others

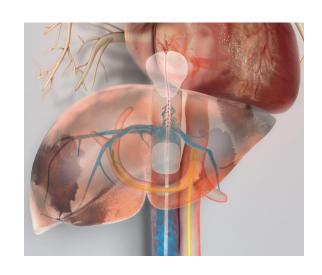






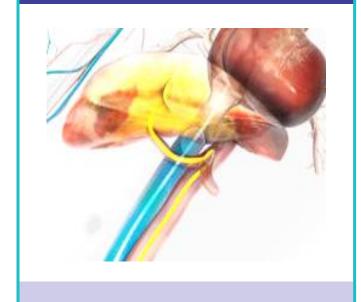


ISOLATION



Isolates hepatic blood flow

SATURATION



A high dose of HEPZATO is delivered to the liver

FILTRATION



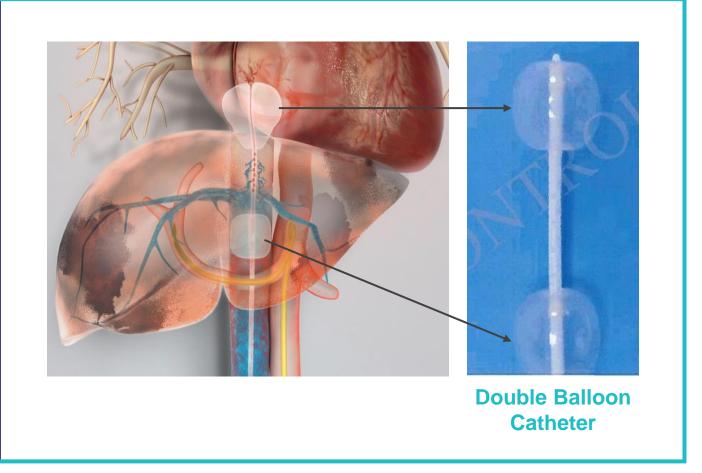
Venous blood from the liver is filtered to reduce melphalan concentration prior to returning the blood to systemic circulation

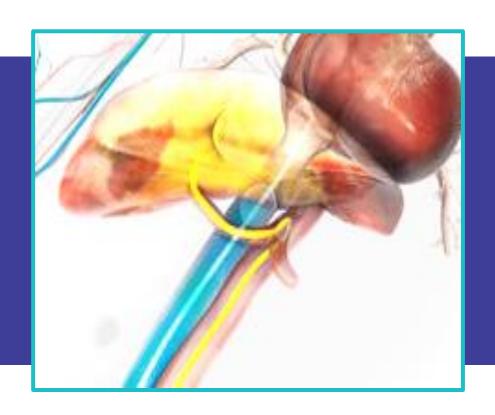
Isolation



The double balloon catheter allows for the isolation of the hepatic venous blood.

Hepatic venous blood flow isolation allows melphalan to be concentrated within the liver.



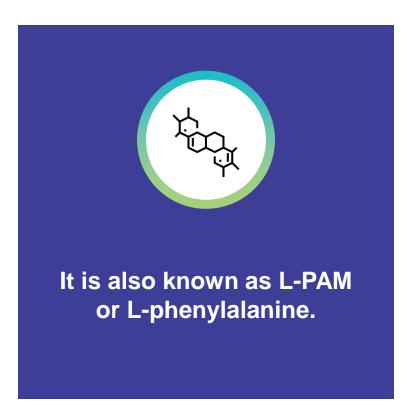


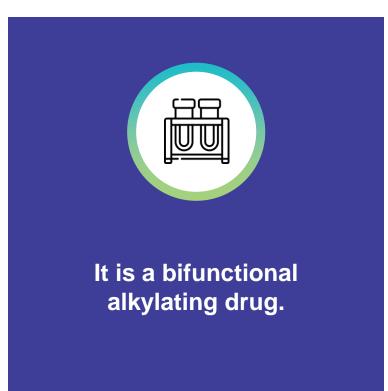
Saturation

Saturation involves a high concentration of melphalan delivery to the liver through the hepatic artery.

HEPZATO (melphalan)



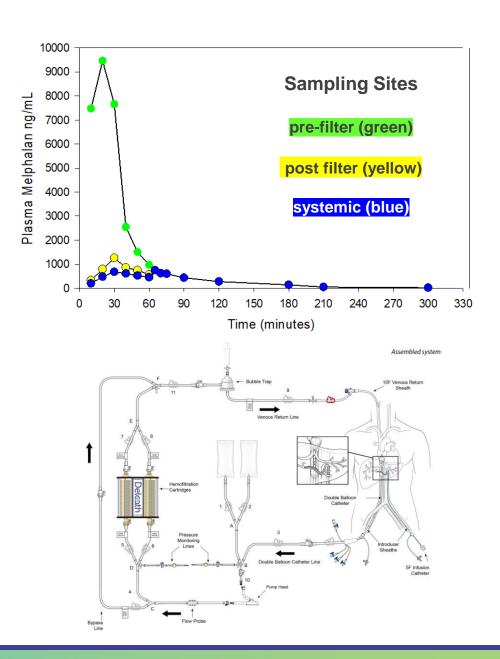




Vahrmeijer AL, van Dierendonck JH, Keizer HJ, Beijnen JH, Tollenaar RA, Pijl ME, et al. Increased local cytostatic drug exposure by isolated hepatic perfusion: a phase I clinical and pharmacologic evaluation of treatment with high dose melphalan in patients with colorectal cancer confined to the liver. Br JCancer. 2000;82(9):1539–46.

Filtration

- The extracorporeal bypass circuit reduces systemic melphalan exposure by ~90%.
- The majority of melphalan does not reach the systemic circulation due to the extracorporeal filtration procedure.



PHP Procedure Team







HEPZATO KIT Clinical Treatment Team



HEPZATO KIT Clinical Treatment Team - Others









Oncologist (Medical Oncologist)





Manage (metastatic) uveal melanoma patients

Monitor and manage chemotherapy toxicities



Communicates

Overall treatment plan and referrals



Refer patient for PHP treatment

Complete medical management of the patient pre- and post-operative care



Coordinates with

Interventional Radiologist, Surgical Oncologist, Anesthesiologist, and other HCPs

Oncologist (Surgical Oncologist)





Monitor and manage chemotherapy toxicities



Communicates

HEPZATO use and risks

HEPZATO KIT procedural risks



May share leadership of procedure with the Interventional Radiologist

Complete medical management of the patient pre- and post-operative care



Coordinates with

Medical Oncologist, Perfusionist, Surgical Oncologist, Anesthesiologist, and other HCPs





Pre-Procedure

- Indications, contraindications, warnings and precautions
- Clinical data summary: Efficacy and safety
- Patient screening and eligibility
- Pre-procedure exam and prescribing pre procedure medications



Procedure

- Procedural related complications
- HEPZATO dosing, preparation and delivery



Post-Procedure

- Mitigating and managing HEPZATO related adverse effects
- Preparing patient for next treatment cycle

Interventional Radiologist





Advanced experience in vascular procedures and liver directed therapies



Communicates

Procedural directives

Vascular anatomy and access

Double balloon inflation



Leads the procedure

Assess vascular anatomy

Access vasculature using sheaths & catheters

Inflate and deflate double balloons



Coordinates with

Perfusionist, Medical/Surgical Oncologist, Anesthesiologist, and other HCPs





Pre-Procedure

- Indications, contraindications, warnings and precautions
- Clinical data summary: Efficacy and safety
- · Patient screening and eligibility
- Identify HEPZATO infusion location based on disease burden and hepatic artery anatomy to ensure adequate drug infusion to the entire liver.
- Pre procedure exam and prescribing pre procedure medications



Procedure

- HEPZATO KIT components and non-KIT required accessories
- HEPZATO dosing, preparation and delivery
- · Vascular access sites and catheterization
- · Hepatic artery mapping and embolization when needed
- Placing hepatic artery infusion catheter
- · Ordering and receiving HEPZATO





Procedure (cont.)

- Introducing and positioning the double balloon catheter (DBC). Technique for balloon expansion, catheter re-position, techniques for assessing and maintaining occlusion, troubleshooting, managing intra-procedural complications and catheter withdrawal.
- · Connecting extracorporeal circuit to patient
- Expanding balloons
- Administering HEPZATO
- Monitoring for hepatic artery patency and/or spasm and methods to relieve spasm.
- · Procedural related complications



Post-Procedure

- Collapsing balloon, disconnecting from extracorporeal circuit and withdrawing catheters
- · Remove sheaths and close access sites
- Monitor for bleeding

Anesthesiologist





Administration and monitoring of general anesthesia

Respiratory and cardiovascular surgical intervention and support



Communicates

Cardiac status – mean arterial pressure, blood pressure

Vasopressor administration

Respiratory status

Post procedure protocol



Management of sedation, analgesia, respiratory and blood pressure/ cardiovascular support

Fluid administration and management

Post procedure management



Coordinates with

Surgical Oncologist, Interventional Radiologist and Perfusionist, Intensivist





Pre-Procedure

- Patient assessment and eligibility for general anesthesia and mechanical ventilation
- Patient screening for cardiac disease including coronary insufficiency, valvular heart disease and left ventricular systolic and diastolic dysfunction



Procedure

- Hemodynamic preparation and monitoring
- Fluid preparation and management
- Anticoagulation preparation and management
- Vasopressor preparation and management
- Preparing for Double Balloon Catheter balloon expansion in the IVC
- Preparing for filters going "online"
- Monitoring and managing extracorporeal filtration system



Post-Procedure

- Reverse coagulopathy
- Monitor and manage hemodynamics
- Maintain and support airway

Perfusionist





Extracorporeal circuit assembly and management



Communicates



When circuit lines are open and closed

Activated clotting time

Pump flow and pressures



Verify KIT type based on IR order

Checks for KIT expiration date

Opens HEPZATO KIT

Records KIT parts serial number before KIT set up

Sets up HEPZATO KIT

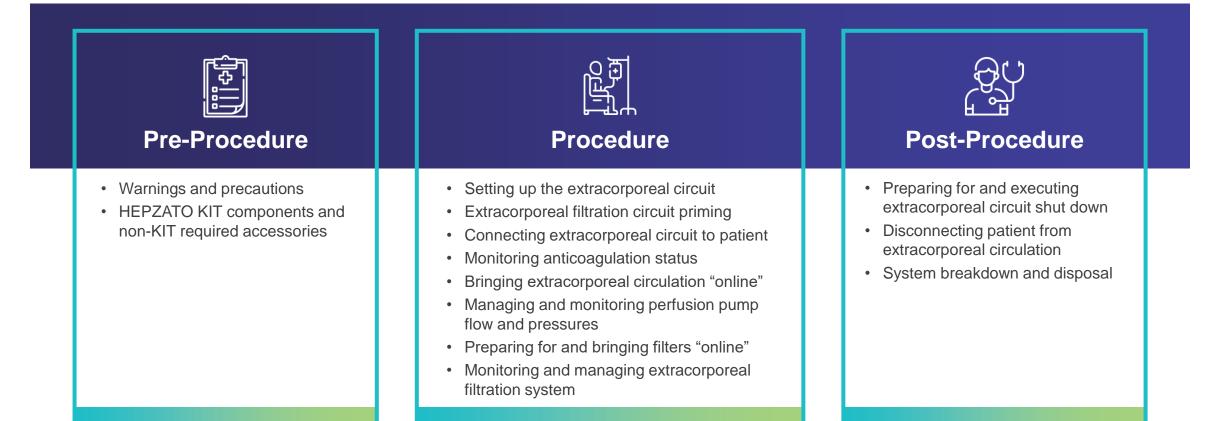
Establish, monitor, controls, and ends extracorporeal pump and veno-venous bypass circuit



Coordinates with

Interventional Radiologist, Anesthesiologist, (Surgical Oncologist), and procedural team







Pharmacist



Preparation of chemotherapeutic agents using national and local safety guidelines



Communicates

During HEPZATO delivery



Reconstitute HEPZATO

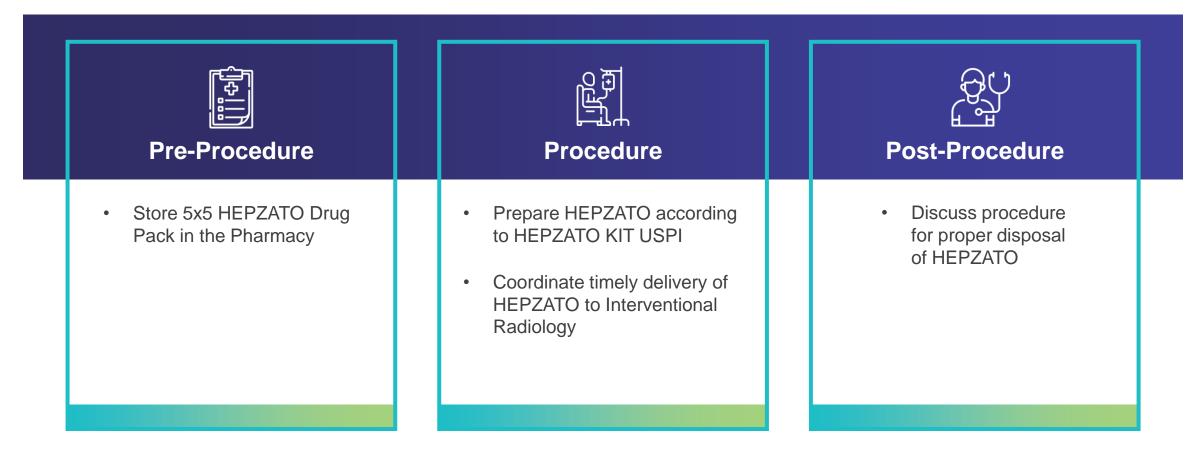
Deliver reconstituted HEPZATO to procedural room



Coordinates with

Medical/Surgical Oncologist, Interventional Radiologist, Registered Nurse, Technicians





Intensivist

Intensivist



Manage critical care patients



Communicates

With the medical or surgical oncologist and Interventional Radiologist



Manages patient immediately postprocedure along with the SO/MO



Coordinates with

Anesthesiologist, Medical/Surgical Oncologist, Interventional Radiologist, Registered Nurse



Pre-Procedure	Procedure	Post-Procedure
 Procedure induced physiological anomalies Procedural related complications 	Complications and adverse reactions	 Manage hematology and administration of blood products Monitor hemodynamic status Monitor for bleeding at access sites and retroperitoneal







MODULE 3
Treatment Planning

Module 3 Purpose and Objectives

The introductory module provides a general overview of HEPZATO KIT and PHP procedure including target patient population and indication, contraindications, general safety and precautions, and clinical efficacy/safety information. Training will prepare HCPs for individual education modules based on specialized roles.



Identify patients who would be suitable candidates for HEPZATO KIT therapy.



Understand importance of pre-procedure imaging.



Know process for preparing patients for HEPZATO KIT procedure.



Understand options for hepatic artery mapping.

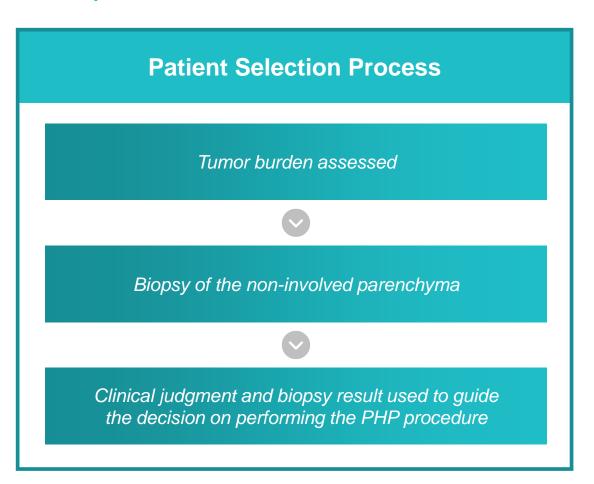
Patient Selection – Characteristics

There are suitable characteristics that reduce procedural/device related risks.



Inclusion

- Patients should have less than 50% liver tumor burden.
- Liver disease must be measured by CT or MRI.
- There can be limited extrahepatic disease at baseline if the life-threatening component of progressive disease is in the liver.
- Patients should have an ECOG score of 0 to 1 at screening.
- Prior chemotherapy, radiotherapy, chemoembolization, or Immunoembolization is allowed with a washout period of 30 days.



HEPZATO KIT Preoperative Patient Assessment

After referral by the Medical Oncologist, a preoperative assessment of the patient is performed by the Anesthesiologist. A Cardiologist, and a Pulmonologist may be consulted for preoperative clearance indicating adequate cardiac and pulmonary function, as per each institution's process or protocol.

A patient may not be eligible for HEPZATO KIT treatment if any of the following are present:

Active coronary artery disease

Severe angina

Recent myocardial infarction

Any congestive heart failure

Significant ventricular arrhythmias

Moderate to severe valvular disease

Unable to withstand high dose vasopressor therapy

Advanced COPD

Unable to withstand mechanical ventilation

Treatment Planning and Preparation, Warnings & Precautions

Treatment Planning and Preparation

Treatment Planning



- No prior liver medical treatments within 4 weeks
- · Screen for prior surgeries that could affect liver vascular anatomy
- Patients should have < 50% tumor burden

Treatment Preparation



- Patients should discontinue chronic anticoagulation therapies
- Patients must discontinue drugs treating hypertension
- Only manipulate intravascular catheters and fluoroscopic guidance
- · Use caution to avoid air embolisms
- · Contents are supplied sterile, inspect prior to use
- All components are single patient use only

Warnings and Precautions



- Peri-procedural complications (including hemorrhage, hepatocellular injury, and thromboembolic events)
- Myelosuppression
- Hypersensitivity Reactions
- GI Adverse Events (nausea, vomiting, abdominal pain, diarrhea)
- Carcinogenic/Mutagenic Effects
- Embryo-fetal toxicity
- Infertility

See Instructions For Use for detailed description of Warnings and Precautions

PHP Procedure Team Communication & Patient Preparation

PHP Procedure Team Communication Tasks

All HCPs should have an understanding and discussion on each expected task required before, during, and after the procedure

The procedural team should have a pre-procedure discussion

The procedural team should review the specific roles of each HCP throughout the procedure

Patient Preparation

The following exams, laboratory, or diagnostic tests should be performed to ensure patient fitness and eligibility such as



Fitness level



Blood tests: full blood count, liver function test, thrombophilia screen, urea, electrolytes



List of current medications and allergies



EKG



Likelihood of pregnancy status



Likelihood of menstruation during PHP procedure window

Pre-Procedure Exams (1)

The preoperative angiograms should include and enable:

Celiac and superior mesenteric artery injection

Evaluate the portal vein for patency during celiac and superior mesenteric arteriography.

The presence of significant portal venous hypertension is a contraindication for Melphalan/HDS

Completely examine the arterial supply to the liver and assess impact on chemotherapy infusion.

Evaluation of portal venous supply, including splenic, superior mesenteric and portal veins – to determine patency and direction of flow

Assessment for variant hepatic arterial and aberrant gastrointestinal (GI) branches – to prevent inadvertent infusion of GI or visceral branches

Pre-Procedure Exams (2)

The preoperative angiograms should include and enable:

Assess liver blood supply and formulate a strategy for catheter placement to ensure adequate drug infusion to the entire liver. If the risk assessment is unfavorable or the anatomic variation is too complex to allow whole liver or sequential lobar catheterization for safe delivery of melphalan, the procedure must not be performed.

A whole liver infusion (dosing) approach will likely require embolization of the gastroduodenal artery, but depends on its origin relative to the side branches of the distal proper hepatic artery. If the infusion catheter tip can be placed sufficiently distally to avoid retrograde reflux into the gastroduodenal artery, then the latter may not need to be embolized.

Depending on vascular anatomy, a sequential lobar approach may be the best administration option which requires splitting the melphalan dose. This will require repositioning of the catheter during the procedure.

Determination of optimal balloon spacing (50mm or 62mm)

Pre-Procedure Exams (3)



Embolization may be performed in order to avoid reflux or infusion into GI or Visceral arteries. Embolized areas may include the following vascular structures:



- Left gastric artery (LGA), Right Gastric Artery (RGA)
- Hepatic variant anatomy (e.g., replaced hepatic artery)
- Aberrant or unusual anatomical variants

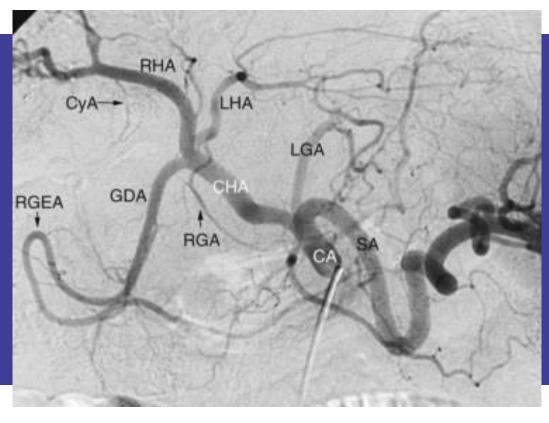
Pre-Operative Angiogram

Thorough search for any variant anatomy

Celiac and superior mesenteric artery mapping

Assessment of the hepatic arterial anatomy

Embolization of certain branches supplying the GI tract



Assess tumor blood supply and formulate a strategy for catheter placement

Select HEPZATO KIT, for balloon spacing, based on patient anatomy

HEPZATO dose planning including whole liver vs lobar delivery

Conventional celiac artery anatomy. CA, celiac axis; LGA, left gastric artery; SA, splenic artery; CHA, common hepatic artery; GDA, gastroduodenal artery; RGA, right gastric artery; RHA, right hepatic artery; LHA, left hepatic artery; CyA, cystic artery; RGEA, right gastroepiploic artery.

Treatment planning options can consist of dosing based on a sequential lobar or whole liver approach.

Typical Schedule of Examinations Prior to PHP Procedure

3 Weeks Prior

- Baseline laboratory medical history and physical examination
- Serum chemistries and electrolytes
- Complete blood count
- Baseline laboratory
- Concomitant disease
- Cardiac and pulmonary evaluation
- Liver function tests
- Prothrombin Time /Partial Thromboplastin Time
- Other medical tests as deemed necessary

2 Weeks Prior

- A baseline CT or MRI is acquired.
- This scan is needed to determine if there has been rapid advancement of the tumor from the time that the diagnostic scan was done.
- The extent of the disease is documented during this scan.
- Procedural risks are assessed including portal hypertension, liver cancer burden, cerebral bleeding risks and histories of surgical procedures.
- Hepatic Artery Mapping (has an optional time point)
- Pre-operative clearance

1 Week Prior

- Patient is screened for blood products.
- Blood Products (Type & Cross)
- Packed RBC, fresh frozen plasma, platelets, cryoprecipitate
- The preoperative hepatic artery mapping study may be performed.
- Pre-op medications prescribed
- Discontinuation of medications
- Hydration instructions
- Pre-op clearance

Procedure Preparation: Prior to Treatment (1)

Hydration May be started the night before the procedure or the day of the procedure

Allopurinol (if tumor burden >25%) As prophylaxis for electrolyte abnormalities, administer 300 mg/day orally 2–3 days before and 2-3 days following perfusion

Proton Pump Inhibitors To prevent gastritis which may occur as a result of regional melphalan absorption during the procedure, administer proton pump inhibitors

Anticoagulation |

- The patient will be systemically anticoagulated with heparin during the procedure.
- Administer heparin to the patient only AFTER placement of the 18F (femoral vein), 10F (jugular vein), and 5F (femoral artery) sheaths
- The patient must be fully heparinized prior to the insertion of the Double Balloon Catheter into the inferior vena cava. Begin with an initial intravenous bolus of heparin at 300 units/kg, dose adjusted to achieve activated clotting time.
- A minimum activated clotting time (ACT) of 400 seconds is necessary with a recommended ACT value > 450 seconds
- Evaluate activated clotting time frequently (approximately every 5 minutes) until adequate anti-coagulation is established (ACT > 400 seconds)
- DO NOT insert double balloon catheter into the patient until ACT values are > 400 seconds.
- Maintain activated clotting time at > 400 seconds throughout the procedure, assessing ACT values every 15 30 minutes depending on the patient's response and by administering intravenous heparin as needed.

Procedure Preparation: Prior to Treatment (2)



Anesthetic Management

Treatment must be administered with patients being monitored and under general anesthesia. Emergency resuscitation equipment must be available during the procedure.



Blood Pressure Control

Expect significant procedure related decrease of blood pressure when the balloons occlude blood return from the inferior vena cava (decreased cardiac inflow) and when the filters are brought into the extracorporeal bypass circuit. The reasons for filter-related hypotension are multifactorial, but hypersensitivity to non-physiological surfaces (inflammatory response), significant reduction in venous return and preload, and possible removal of catecholamines by the filters may play a role. To aid blood pressure maintenance for extracorporeal bypass, the following actions are recommended per institutional practice:

- Pre-operative hydration and intra-procedural administration of colloids and crystalloids.
- Vasopressor use in accordance with institutional practices to elevate mean arterial pressure to a target >65 mmHg
- Blood pressure must be constantly monitored throughout the procedure and maintained at levels required for adequate perfusion of critical end-organs (i.e., >65 mmHg).

Pre-Procedure Infusion Plan

Coordination with the Pharmacy

Prior to set up, provide prenotification to the hospital pharmacy to be ready to prepare HEPZATO. An actual request for drug preparation and delivery should be timed so that the start of the infusion of HEPZATO is within thirty minutes of preparation.

The filters should not be brought online until HEPZATO is in the procedure room. This minimizes filtration duration and associated complications.

Drug administration should be completed within 60 minutes of the start of preparation.



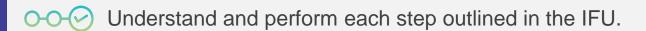


MODULE 4
Perfusion Circuit

Module 4 Objectives



Module 4 describes the responsibilities of the perfusionist in assembling the EFC and collaboration with the IR when connecting the patient to the extracorporeal circuit. HCPs will gain an understanding of the perfusion circuit components, setting up the system, managing the fluid path, connecting the system to the patient, bringing the filters online, troubleshooting flow related issues, and disconnecting the patient from the EFC.





Interpret extracorporeal pump flow metrics.



Support proper anticoagulation in collaboration with the anesthesiologist.



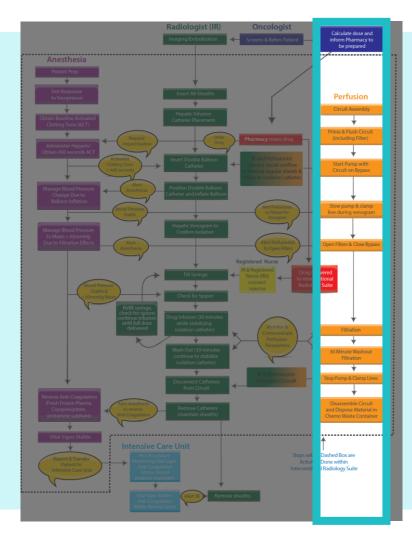
Identify EFC issues and know how to resolve these problems.



Communication between the Perfusionist and the **other team members** is of **critical importance**.

Perfusionist Procedural Tasks





This module describes the steps the Perfusionist takes during the procedure



Circuit Assembly

Prime & Flush Circuit (including Filter)

Start Pump with Circuit on Bypass

Slow Pump & Clamp Line During Venogram

Open Filters & Close Bypass

Filtration

30 Minute Washout Filtration

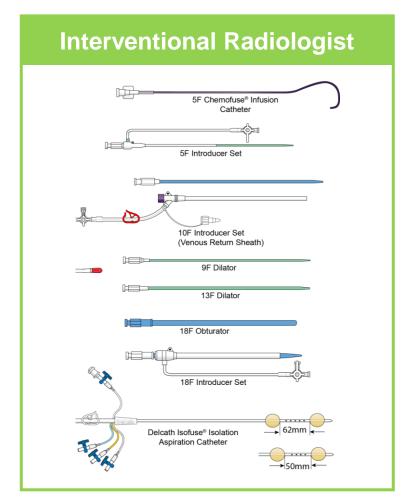
Stop Pump & Close Lines

Disassemble Circuit and Dispose Material in Chemo Waste Container

HEPZATO KIT Components



Perfusionist

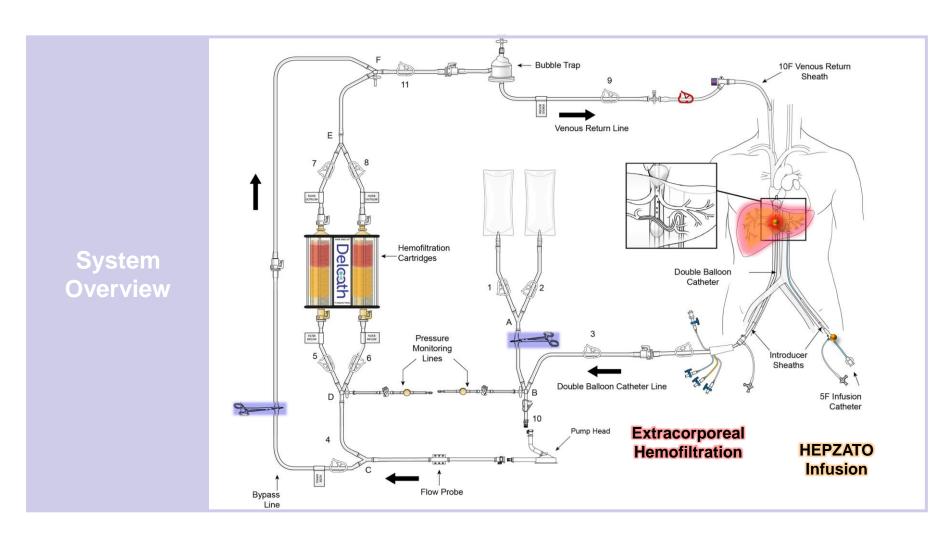


Only components provided in the Hepatic Delivery System or specified by Delcath in the Instructions For Use are to be used to create the circuit.

See IFU for full system description, including alphanumeric descriptions







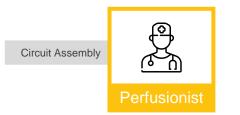
NOTE

Numbers 1-11 denote clamps used to control flow

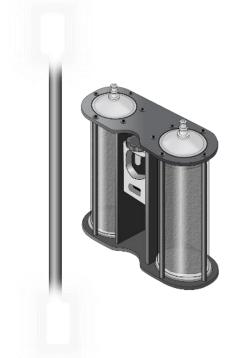
Letters A-F denote Y-connectors

See IFU for full system description, including alphanumeric descriptions

1. Assemble the Extracorporeal Filtration Circuit



- Prepare 9 liters of Heparinized Normal Saline
- Remove Hemofiltration
 Dual Filter Cartridge from
 sterile pouch
- Using guidance from the "THIS END UP" label on the faceplate, attach the filter to the intravenous pole using the built-in pole mount clamp.



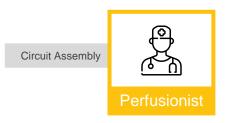


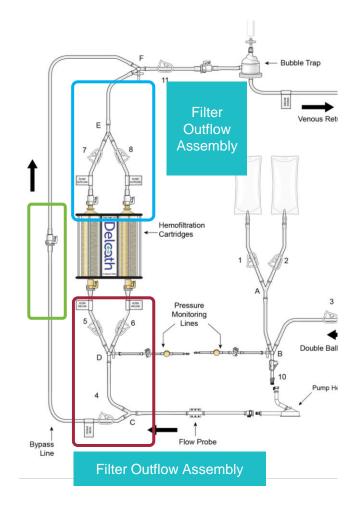


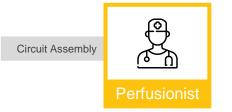


Filter & Circuit Assembly

- 1. Open the circuit tray and remove the components in the pouches. Set aside for later assembly.
- 2. Remove the "Filter Inflow Assembly" and "Pump Head Assembly" sections from the circuit tray, connect Filter Inflow Assembly to Pump Head outlet and DBC Assembly to Pump Head inlet. Place pump head on the pump drive motor, and insert flow probe into flow transducer.
- Connect the Inflow lines (circled in red in the image to the right) to the cartridge inlet connectors located on the bottom of the dual filter.
- 4. To complete the assembly, push the quick connector couplings together, as shown (circled in red; male to female) until an audible "click" is heard to verify connection.
- 5. Remove the "Filter Outflow Assembly" section from its sterile pouch.
- 6. Connect the two outflow lines to the filter cartridge outlet connectors located on the top of the dual filter using the quick connector couplings (circled in blue; push and click).
- 7. Assemble the two ends of the **Bypass line** by pushing the quick connectors together (circled in green; push and click).

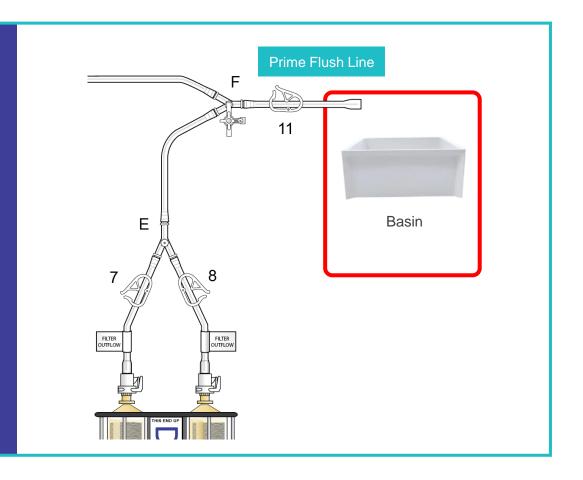






Prime Flush Line Assembly

- Remove the "Prime/Flush Line" from its sterile pouch.
- Attach the prime/flush line to the filter outflow assembly using the quick connect coupling located distal to "Y"-connector at F (blue arrow).
- Place the open end of the "Prime/Flush Line" into the basin for collecting the flushed effluent during filter hydration (circled in red).



Perfusionist

Extracorporeal Filter Circuit Setup

Stopcock Attachment Locations

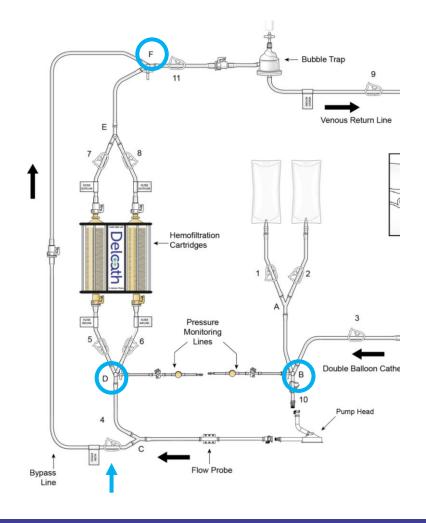
Attach the supplied stopcocks as identified by the blue circles to ports:

"B"
(pre-pump head
pressure/suction)

"D" (pre-filter pressure)

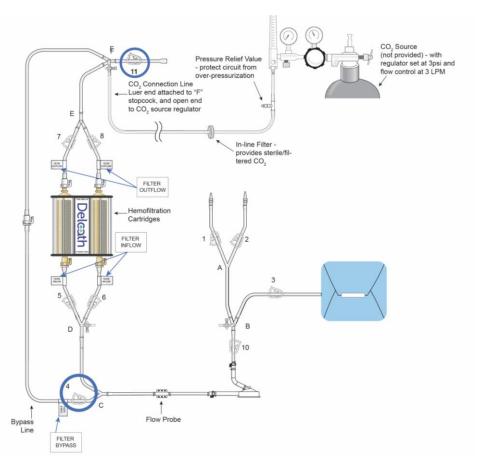
"F" (outlet) Y-connector

- Ensure all stopcocks are in the closed position (lever closed to the perfusion circuit).
- **3** Verify "Bypass Line" clamp 4 is open (Blue Arrow).





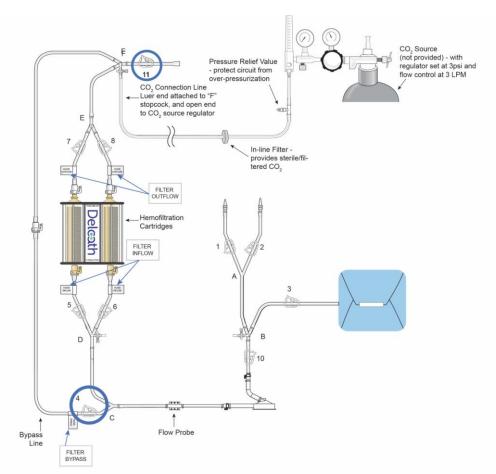
CO₂ Priming Connections



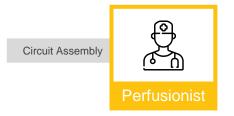
- 1. Close outlet clamp 11.
- 2. Attach the CO² connection line to the stopcock "F" and open the stopcock.
- 3. Connect the open end of the CO² connection line to the CO² source, and set the CO² source regulator to 3psi (approximately 3.0 liters per minute).
- 4. Start the CO² gas flow and allow the CO² to flow through the hemofiltration circuit.
- 5. Adjust CO² regulator to maintain 3psi (as necessary).



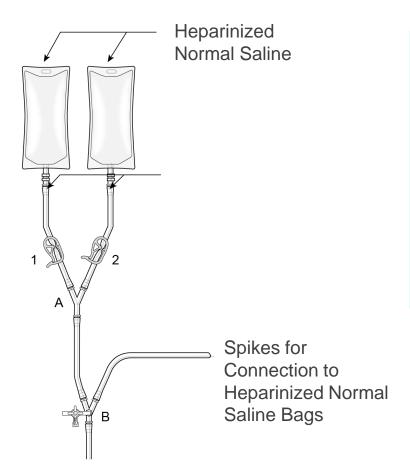
CO₂ Priming Connections



- 6. Verify CO² flow through the circuit.
- 7. Close bypass clamp (4) after approximately 1 minute to ensure flow through the hemofiltration cartridges.
- 8. Allow CO² to flow through the cartridges (after closing clamp 4) for at least 5 minutes.
- 9. Leave saline clamps 1 and 2 closed. Close double balloon catheter line, clamp 3. Close filter inlet clamps 5 and 6. Then close filter outlet clamps (7, 8) to lock CO² within the circuit.
- 10. Stop the CO² flow and close the stopcock "F" and disconnect the CO² connection line and discard.



Heparinized Saline Bags Connections





Hang two bags of the heparinized sterile normal saline and connect to circuit by using the spikes to allow for gravity priming of circuit components.

IMPORTANT: Use strict aseptic technique while spiking the heparinized normal saline bags.

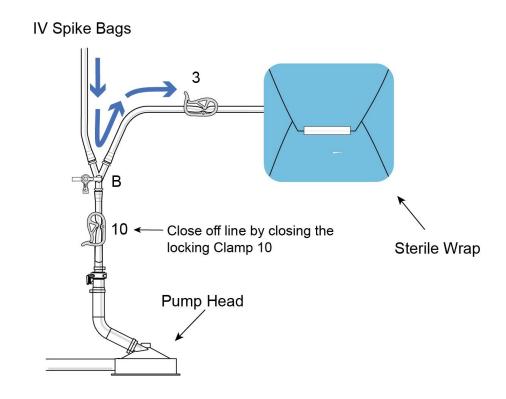
Ensure there are two (2) liters of normal saline available for later use

Circuit Assembly

Extracorporeal Filter Circuit Setup

Prime Double Balloon Catheter Line

- Close pre-pump clamp 10.
- Confirm that stopcock "B" is in the closed position (the lever is pointed toward the circuit tubing to the right).
- Open double balloon catheter line clamp 3.
- Open saline line (clamp 1 or 2), to allow heparinized normal saline to prime line only up to clamp 3, blue arrows demonstrating flow direction. Do not allow excess heparinized normal saline to fill sterile wrap.
- Close clamp 3.



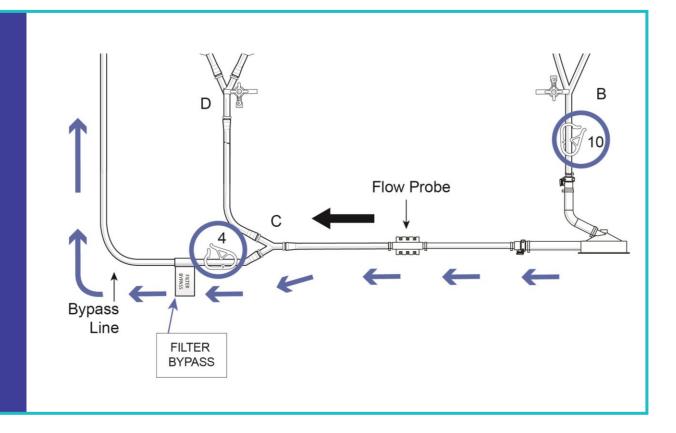
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Extracorporeal Filter Circuit Setup

Circuit Assembly Perfusionist

Prime Bypass Line

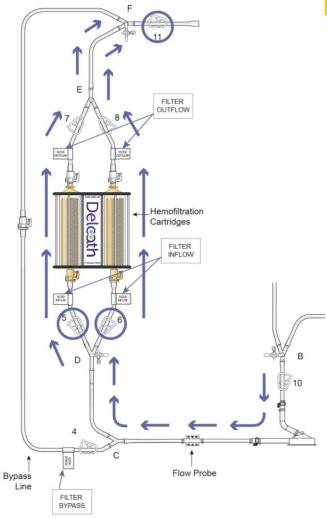
- Inspect that stopcock "D" is in the closed position (the lever is pointed toward the circuit tubing to the left).
- Prime the pump head and bypass lines by opening clamps 10 and 4.
- Close clamp 4.





Prime Filter Cartridges

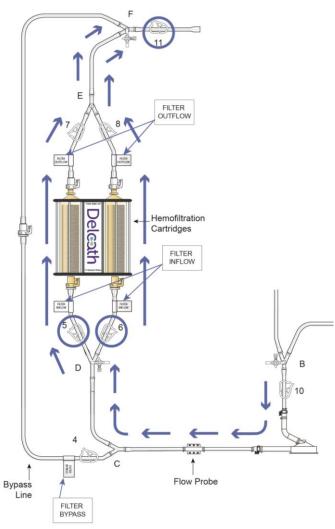
- 1. Inspect that stopcock "F" is in the closed position (the lever is pointed upward).
- 2. Open filter inlet clamps (5, 6).
- 3. Open filter outlet clamps (7, 8).
- 4. Open circuit outlet clamp (11).
- 5. Adjust the flow of heparinized normal saline into the filter to a suggested starting rate of 0.1 liters per minute (increasing slowly up to 0.5 liters per minute when the filters are completely filled). Note: Hemostats (forceps) are required to adjust flow rate if using gravity.
- 6. Note that filter cartridges will have a mottled appearance indicating the presence of gas bubbles.





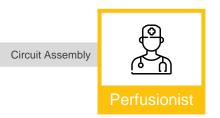
Prime Filter Cartridges

- 7. Allow heparinized normal saline to flow through the filters and out the "Prime/Flush Line" for approximately 6-10 minutes or until the filter appears gas free (solid black).
- 8. Once all gas appears to have been displaced, gently roll the cartridges between palm of hands to encourage trapped gas bubbles to rise. CAUTION: Do not use excessive force when rolling the plastic housing.
- 9. Inspect the entire cartridge for trapped gas by turning the cartridge within housing to visualize the entire filter. If there are gas bubbles gently roll the cartridges to free the trapped gas.
- 10. When Filter Cartridges are gas free, flush with an additional six (6) liters of heparinized normal saline (3 L/cartridge).
- 11. Close filter clamps 5, 6, 7, 8 and outlet clamp 11.



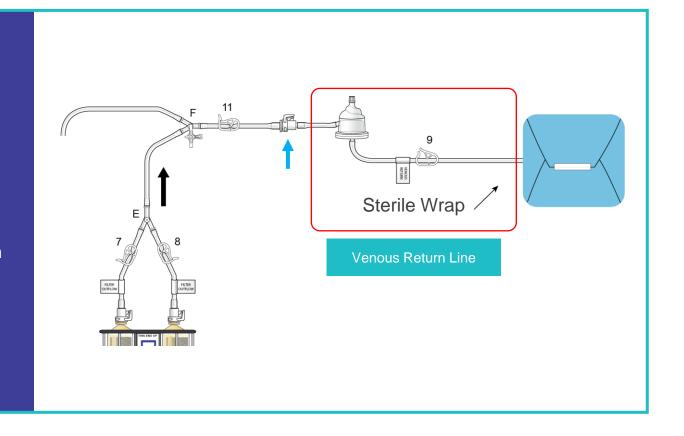
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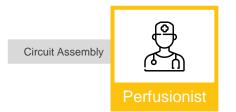
Extracorporeal Filter Circuit Setup



Prime Venous Return Line and Bubble Trap (1)

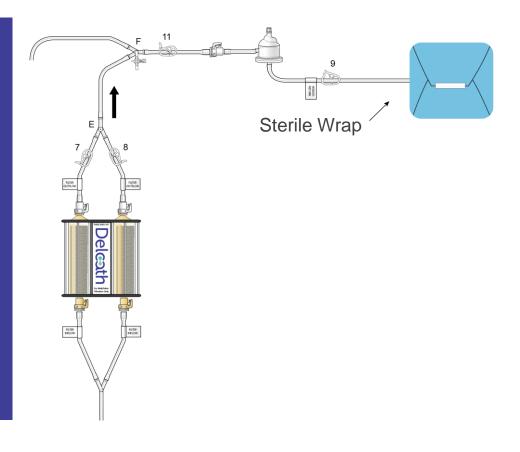
- Disconnect and dispose of the "Prime/Flush Line", by pressing in the latch located on the female quick connect coupling and pulling it apart.
- Open the Venous Return sterile pouch and remove the venous return line and built-in bubble trap.
- Attach the female to the male quick connector (push and click) located (blue arrow) by outlet clamp (11).
- Position the bubble trap in the bubble trap holder higher than filter cartridges.

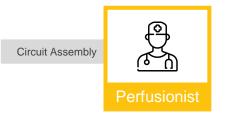




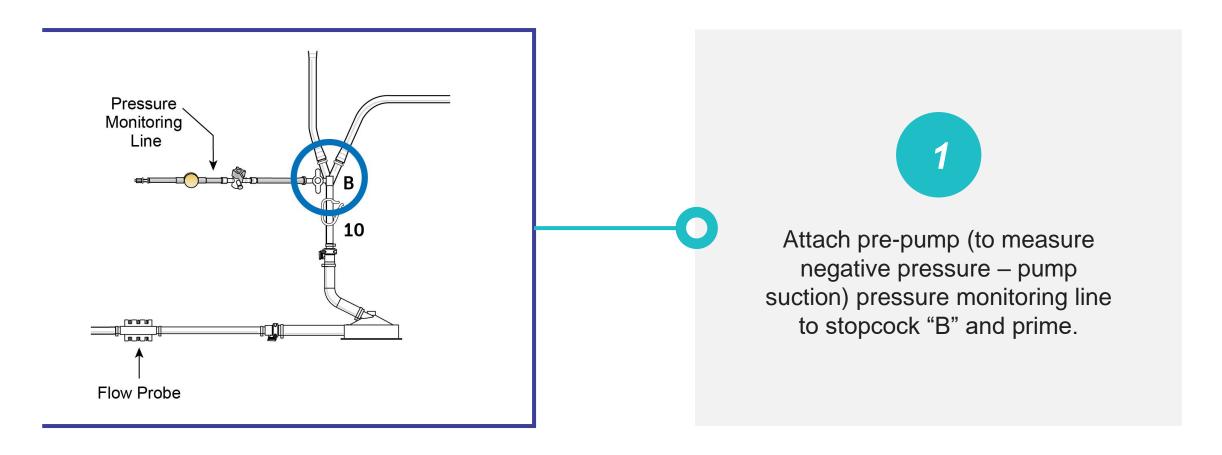
Prime Venous Return Line and Bubble Trap (2)

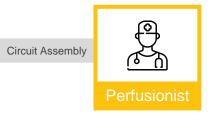
- Attach stopcock to bubble trap and use syringe to aspirate air, as necessary.
- Prime venous return line and bubble trap by opening clamps 4, 11 and 9.
- Prime up to clamp 9. Do not allow saline to enter blue sterile pack.
- Close clamp 9 once venous return line and bubble trap are primed up to clamp 9.





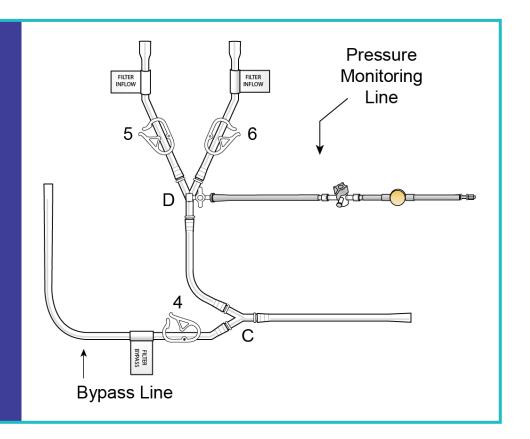
Pre-Pump Pressure Line Attachment & Priming (1)





Pre-Pump Pressure Line Attachment & Priming (2)

- Attach pre-filter (to measure positive pressure pre-filter) pressure monitoring line to stopcock "D" and prime.
- Attach the pressure monitoring lines to the P1 and P2 ports on the rear of the Medtronic Bio-Console 560 Speed Controller System.
- Zero the pressure transducers (refer to Medtronic Bio-Console 560 System Manual for details).
- Coiled pressure monitoring lines are included for use with DLP Pressure Display Boxes, as necessary.



E

Extracorporeal Filter Circuit Setup



Circuit Assembly

Pressure Test Circuits

- Pressure test circuit by slowly ramping up the pump head speed (RPM) until a pressure reading of 300 mmHg is achieved on the pressure transducer attached to the line on Y-connector "D" (pre-filter).
- 2 Visually inspect all connections and cartridges to ensure no leaks are present.
- **3** Turn off pump and close cartridge inlet (5, 6) and outlet (7, 8) clamps. Ensure bypass line clamp (4) is open.
- 4 Ensure there are two liters of normal saline available for later use.
- 5 Warning: Ensure that all air is purged from the system prior to use in order to avoid an air embolism.

ECF System is now primed, hydrated, de-bubbled, and ready for use



Perfusionist Procedural Awareness

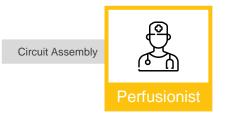
Pump Flow Rate

The flow rate of the pump should be maintained between 0.4 – 0.8 L/min

Pre pump pressure (suction side) should not be more negative than -250 mmHg, as lower pressures indicate possible catheter collapse or kink.

Pre-cartridge pressures (pre-filter) should not exceed 200 mmHg, as higher pressures indicate increasing filter resistance potentially due to thrombus or a kinked return line. Check filters to assure free flow and return line for kinks.





Before Veno-Venous Bypass



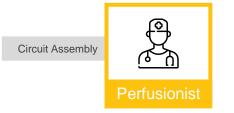
Nurse/Tech Communicates: HEPZATO has been delivered

Prior to opening the veno-venous bypass circuit:

HEPZATO should be fully reconstituted and prepared for injection.

2 The activated clotting time should be > 400 seconds.

Vasopressor agents should be prepared and ready for injection.

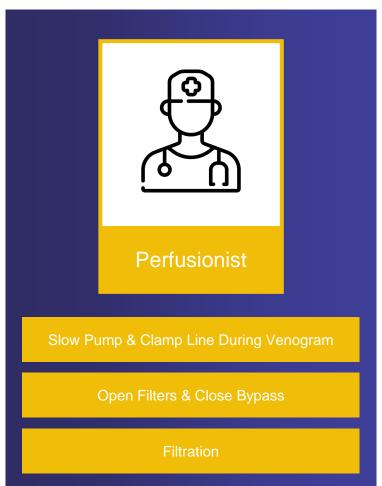


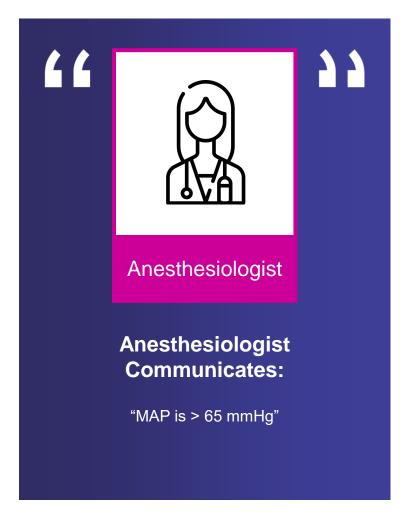
Veno-Venous Bypass

Pump Speed	Extracorporeal Filtration Circuit Clamps	Blood pressure
Starting the Medtronic pump at a low speed to allow blood flow from the double balloon catheter into perfusion circuit.	It is important to open all of the clamps EXCEPT for the filter inlet and outlet clamps, which are clamps 5-8. During veno-venous bypass, blood will bypass the filters and be returned to the internal jugular vein.	Blood pressure should be carefully monitored, and the mean arterial pressure should be above 65 mmHg.

HCP Communications Before Bringing Filter Cartridges Online







Bringing Hemofiltration Cartridges Online

Open Filters and Close Bypass









Leaving the bypass line open, open clamps on the left cartridge (5 and 7) and allow blood to displace the heparinized normal saline into the patient.

After the heparinized normal saline is in the left cartridge and its lines is fully replaced with blood, wait approximately 30 seconds and open clamps on the right cartridge (clamps 6 and 8), while keeping the bypass line open.

Once the heparinized normal saline in the right cartridge and its lines is fully replaced with blood, wait approximately 30 seconds and then close the bypass line by securely closing clamp 4.

Add a reusable tube clamp as a redundant bypass closure mechanism high on the bypass line in clear view of the team.







Perfusionist Communicates:

"Filter 1 is open"

"Filter 2 is open"

"Bypass line is closed?"

Anesthesiologist Communicates:

"Blood Pressure is..."

"MAP is controlled"

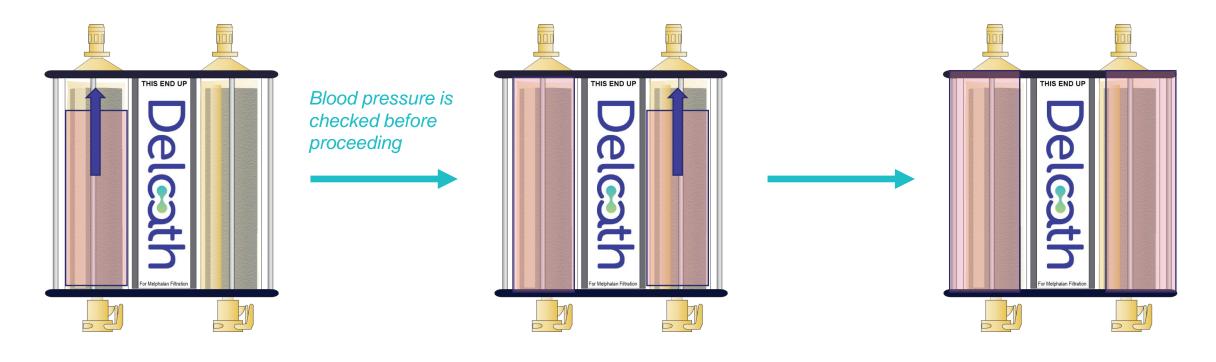
Start Pump with Circuit on Bypass

Slow pump & clamp line during venogram



Hemofiltration Cartridges Method

Filters brought online sequentially to limit large blood pressure reduction



Left cartridge is opened and charged first

Right cartridge is then opened and charged

Complete Filtration Established



The tube clamp can be used to close the bypass line, at the area close to the upper Y connector and in easy view of the procedural team.

This will ensure that every HCP can see that the bypass line is closed during filtration.

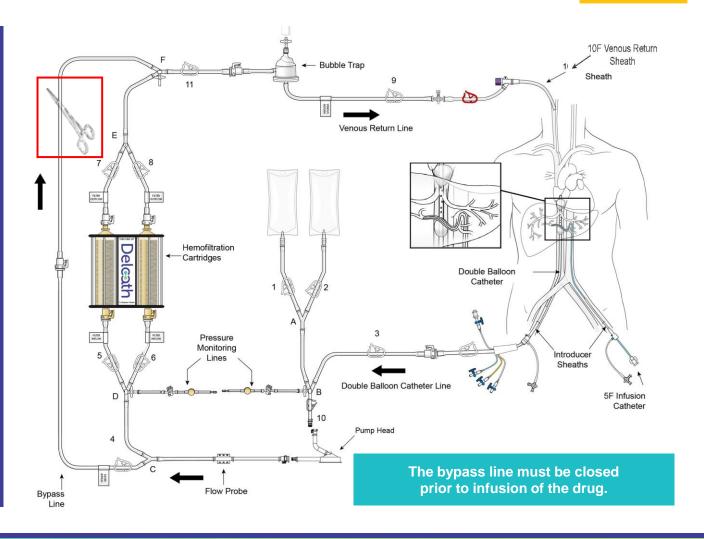




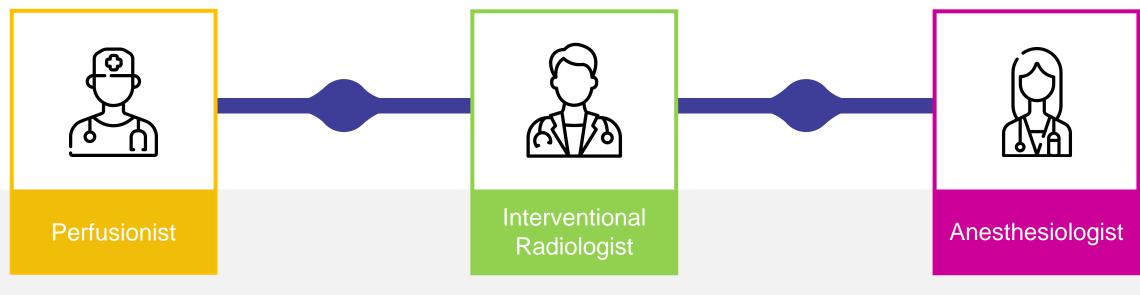


Anesthesiologist Communicates:

"Blood Pressure is..."



PHP Procedure Monitored Events & PHP Team Member Roles



Monitor Bubble-trap for entrapped air

Check for leaks from any part of the circuit

Monitor blood flow rate

Maintain DBC position

Spasm check

Check for DBC leaks

Positioning infusion catheter

Monitor systolic, diastolic, and mean arterial blood pressure

Monitor heart rate and vital signs

Correct as needed



Establishing Hemofiltration

Blood Flow During Hemofiltration



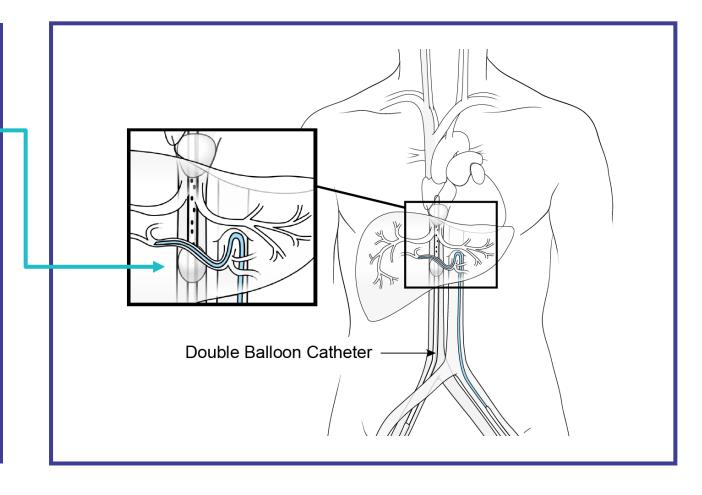
Venous blood is aspirated from the central lumen through the fenestrations in the Isofuse Catheter



This blood flows through the Isofuse Catheter to the pump through the Bypass Line



Blood is returned to the patient through the venous return sheath



30 Minute Washout Filtration



Drug Infusion

Hemofiltration is maintained for

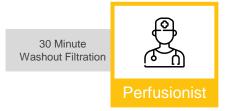


First 30 minutes:
Drug infusion phase

Last 30 minutes: Washout period



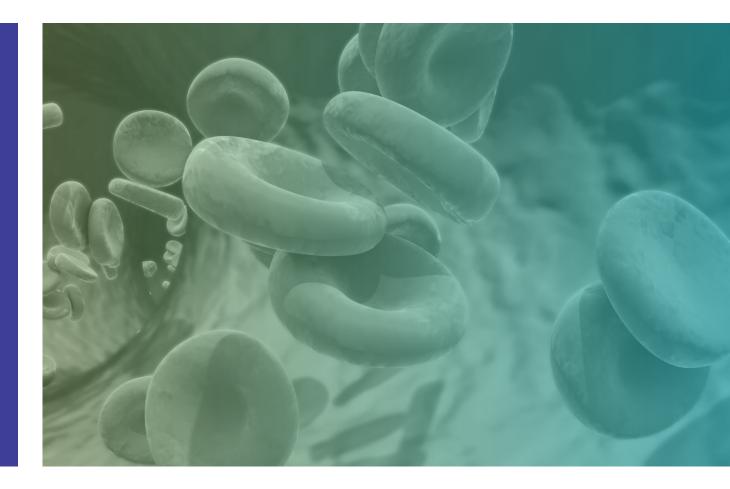
- Helps to maintain hemofiltration
- Monitors blood flow rate
- Monitors Pump Pressures
- Checks for circuit leaks
- Records ACT



Blood Return

To return blood to the patient the Interventional Radiologist instructs the Perfusionist to open clamps for saline flush and this pushes the blood through filter and back into patient's internal jugular vein.

The effectiveness of the return of blood to a patient is dependent on the central venous pressure and the amount of fluid given.







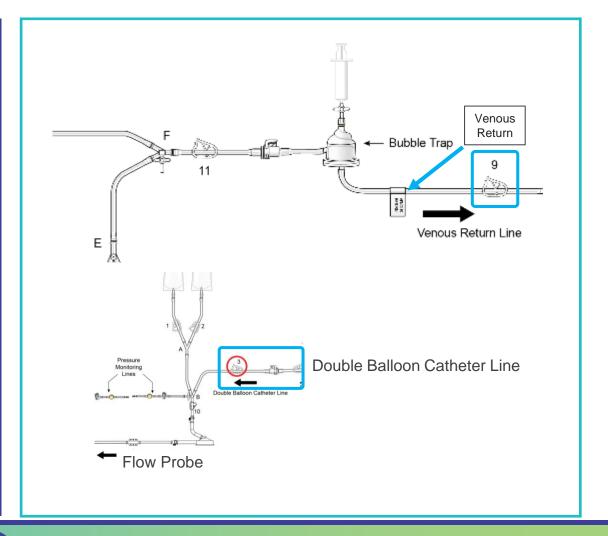
At the end of the 30-minute wash-out period:

- 1 IR fully collapses the caudal balloon.
- 2 IR fully collapses the cephalad balloon.
- **3** Perfusionist discontinues filtration by:

Reducing the pump RPM to 1000

Closing clamps 3 and 9

Stopping flow by turning off the pump.



Disassemble Circuit and Dispose Material in Chemo Waste Container





Perfusionist

Stops all ECF flow

Opens bypass line

Assists with blood return

Appropriately disposes all components



Interventional Radiologist

Collapses DBC

Disconnects DBC

Disconnects internal jugular introducer from the ECF



Anesthesiologist

Vasopressor weaned/discontinued

Normalizes coagulation

Post procedural medications



MODULE 5

Drug Preparation and Injection

Module 5 Objectives

Module 5 describes the responsibilities of the pharmacists and interventional radiology nurse/technician who are responsible for the drug preparation and delivery system preparation, respectively.



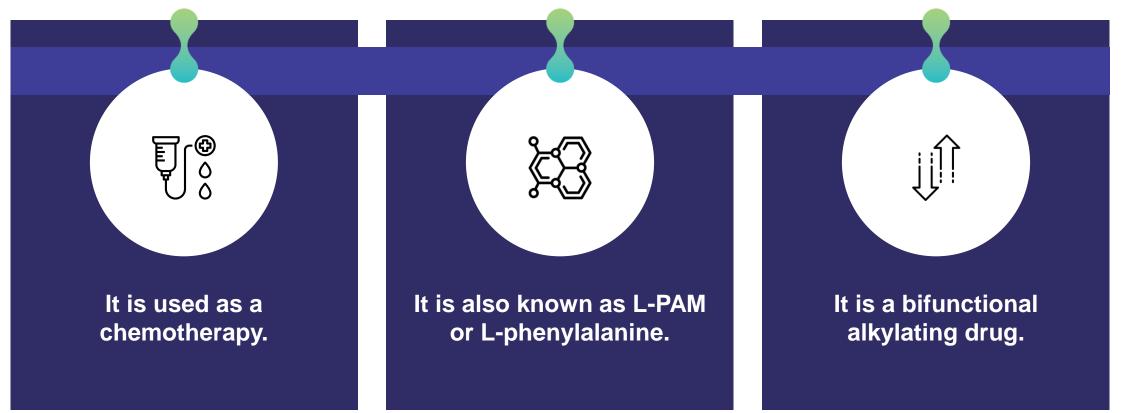
Understand how to reconstitute, store and transport HEPZATO.



Coordinate timing of the prepared HEPZATO so delivery occurs as soon as possible to limit melphalan degradation and procedure time for the patient.



HEPZATO (melphalan) Injection



Vahrmeijer AL, van Dierendonck JH, Keizer HJ, Beijnen JH, Tollenaar RA, Pijl ME, et al. Increased local cytostatic drug exposure by isolated hepatic perfusion: a phase I clinical and pharmacologic evaluation of treatment with high dose melphalan in patients with colorectal cancer confined to the liver. Br JCancer. 2000;82(9):1539–46.

HEPZATO (melphalan) for Injection



- · Wear gloves in preparation and handling.
- If skin or mucosa contact is made, immediately wash with soap and water.
- Follow institutional guidelines and protocols.



Each vial of HEPZATO for injection contains melphalan equivalent to 50 mg melphalan and 20 mg povidone



0.9% Sodium Chloride Injection is used as the approved secondary diluent

Calculation of Ideal Body Weight for HEPZATO Dosing

	HEIGHT	IDEAL BODY WEIGHT
Men ♂	≥ 152 cm	52 kg + (0.75 kg/cm of height greater than 152 cm)
	< 152 cm	52 kg - (0.75 kg/cm of height less than 152 cm)
Women Q	≥ 152 cm	49 kg + (0.67 kg/cm of height greater than 152 cm)
	< 152 cm	49 kg – (0.67 kg/cm of height less than 152 cm)

The HEPZATO dose may be calculated 1 to 2 days before the treatment by the Medical Oncologist.

TOTAL HEPZATO DOSE = CALCULATED IDEAL BODY WEIGHT X 3 MG

NOTE: Total dose not to exceed 220 mg.

HEPZATO Preparation



Prepared by Pharmacy per physician's prescription

Reconstitute each HEPZATO vial with 10 mL of supplied diluent



Doses up to 110 mg

dilute reconstituted
HEPZATO in 250 mL 0.9%
sodium chloride injection

Doses between 111 mg – 220 mg

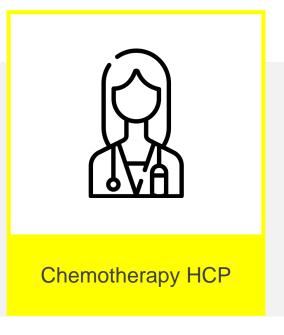
dilute reconstituted
HEPZATO in 500 mL 0.9%
sodium chloride injection

The 250 mL and 500 mL solutions will require multiple injection cycles

HEPZATO Preparation



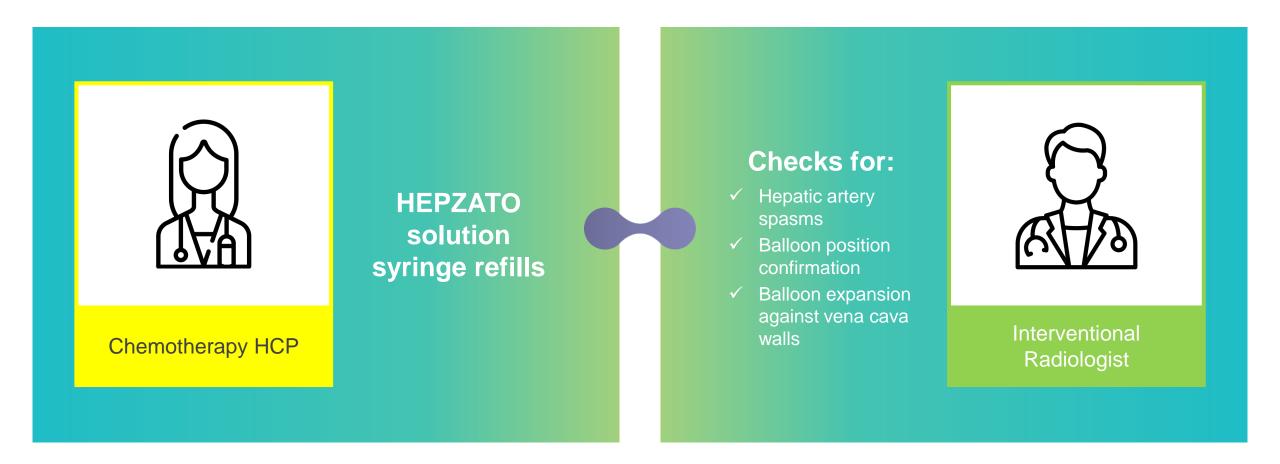
Visual inspection for particulates in HEPZATO solution





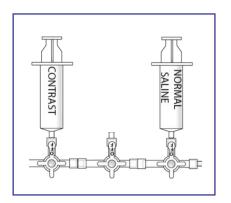
IF PARTICULATES ARE OBSERVED DO NOT USE

HEPZATO Infusion Process



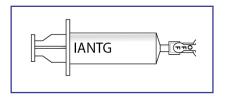
HEPZATO Administration Injection Line Setup

Syringes



Undiluted iodinated contrast agent is attached to the injection line, and it is used for checking hepatic artery spasm via CT.

Nitroglycerin should be kept in the procedural room, and it must be administered intra-arterially to alleviate spasms.



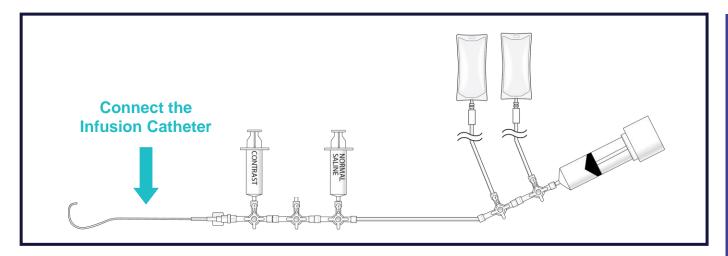
Normal saline is attached to the injection line, and it is drawn into the saline syringe for priming and flushing the hepatic arterial infusion line during the procedure.

Syringe	Volume and Concentration to Prepare in Syringe
Contrast	20 mL of undiluted iodinated contrast agent
IANTG	5 mL of diluted nitroglycerin 100 mcg/mL solution
Saline	20 mL of 0.9% Sodium Chloride injection

HEPZATO Administration Injection Line Setup

Catheter Connection

Note: this connection is done after drug arrives in the room.



- Connect the Infusion Catheter (5F catheter or a microcatheter coaxially introduced through the 5F catheter).
- Maintain catheter patency by hospital catheter infusion protocols (e.g., infuse heparinized saline: the concentration of heparin should be 1000 units per 500 mL of normal saline).

Items needed but not included in HDS

- (1) Medrad 150 mL syringe
- (2) Intravenous Administration Sets
- (3) 48" Injector Lines
- (5) 3-way stopcocks
- (3) 20 mL syringes



Arterial Catheter



HEPZATO is administered by infusion via a 5F arterial catheter or a Delcath-qualified microcatheter.



It is introduced coaxially through the 5F catheter into the proper hepatic artery.



At the discretion of the Interventional Radiologist, a microcatheter may be used when selective tip placement is preferred for the drug infusion.



Delcath has qualified three different microcatheters for use with the HEPZATO KIT.

Drug Injector System Setup

150 mL syringe installed into the drug injector

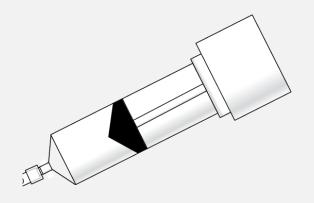
Injector flow rate set to 25 mL/minute OR 0.4 mL/second

Injection volume entered 100 mL per injection

Requires multiple injection cycles



HEPZATO infusion must be completely administered within 30 minutes



HEPZATO Drug Injector & Infusion Timing

Legend Key







Refill Syringe. Check for Balloon/Catheter Placement & Spasm

500 mL Total Infusion (25 mL/min)



250 mL Total Infusion (25 mL/min)



HEPZATO Delivery Parameters & Precautions

DELIVERY PARAMETERS

500 mL

5 infusions of 100 mL at 25mL/min

250 mL

2 infusions of 100 mL + infusion of 50 mL at 25 mL/min

DELIVERY PRECAUTIONS



Immediately stop the procedure if leakage is detected outside of the isolated region and cannot be corrected.



Once the infusion of HEPZATO has started, Do NOT deflate balloons unless administration of drug has been stopped and a full washout cycle (30 minutes) has been completed.







MODULE 6

Procedure/Treatment Day

Module 6 Objectives





Describe required pre procedure imaging, laboratory testing, and medications.



Understand team coordination and intraprocedural communications.



Know procedure day medication, supplies and blood product needs.

Typical Schedule of Examinations Prior to PHP Procedure

3 Weeks Prior

- Baseline laboratory medical history and physical examination
- Serum chemistries and electrolytes
- Complete blood count
- Baseline laboratory
- Concomitant disease
- Cardiac and pulmonary evaluation
- Liver function tests
- Prothrombin Time /Partial Thromboplastin Time
- Other medical tests as deemed necessary

2 Weeks Prior

- A baseline CT or MRI is acquired.
- This scan is needed to determine if there has been rapid advancement of the tumor from the time that the diagnostic scan was done.
- The extent of the disease is documented during this scan.
- Procedural risks are assessed including portal hypertension, liver cancer burden, cerebral bleeding risks and histories of surgical procedures.
- Hepatic Artery Mapping (has an optional time point)
- Pre-operative clearance

1 Week Prior

- Patient is screened for blood products.
- Blood Products (Type & Cross)
- Packed RBC, fresh frozen plasma, platelets, cryoprecipitate
- The preoperative hepatic artery mapping study may be performed.
- Pre-op medications prescribed
- Discontinuation of medications
- Hydration instructions
- Pre-op clearance

HEPZATO KIT Pre-Operative Activities

Ensure timely scheduling of preoperative activities

Imaging studies

Laboratory testing

Medication requirements including new prescriptions



Procedural room reservation

Hospital bed reservation

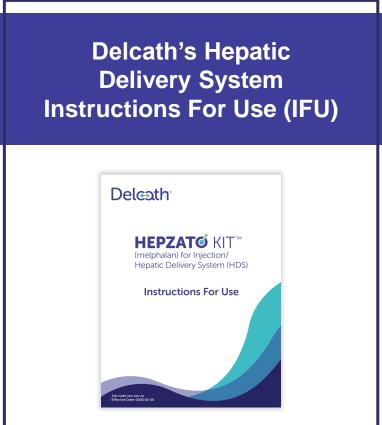
Appointments with:

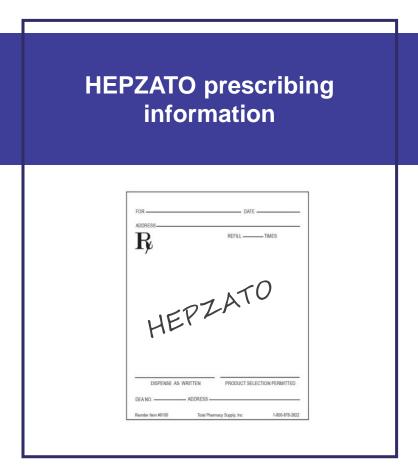
Medical Oncologist
Interventional Radiologist
Anesthesiologist

HEPZATO KIT Treatment Day Guidance

All medications and supportive measures must be determined and administered in accordance with:







The Perfusionist verifies the following:

- The expiration date, reference number, and lot number of the KIT.
- 2 The contents of the HEPZATO KIT.
- 3 Sets up the extracorporeal filtration circuit.
- A full tank of CO₂ is in the procedural room.
- 5 The ACT machine is in the procedural room.



Perfusionist

The Interventional Radiologist verifies the following:

- The correct HEPZATO KIT (balloon spacing 50mm or 62mm) is in the procedure room.
- The time that HEPZATO will be delivered by the pharmacy.
- 3 Any recent patient lab results which are pertinent to the procedure.
- 4 The treatment start time.
- That all the medical materials needed for the procedure are in the operating room.
- 6 That all PHP Procedure team members are in attendance.



Interventional Radiologist

The Anesthesiologist verifies the following:

The anesthesia medications are in the procedural room.

The vasopressor pump or line is set up.

There is patient access for delivery of medications.



Anesthesiologist

The Chemotherapy HCP verifies the following:

1 The drug injector is in the room.

2 The drug injector is set up.

The components that are not included in the HEPZATO KIT are available and organized on the Interventional Radiology suite's tables.



Chemotherapy HCP

Patient Treatment Day Preparation

The patient is:



- Provided with details on forbidden medications (oral anticoagulants, ACE inhibitors, calcium channel blockers, alpha-1adrenergic blockers, thrombin inhibitors, aspirin, NSAIDs)
- Provided with details about medications that will be administered during the procedure
- Administered pre-procedure medications



The patient must be properly hydrated before the procedure.

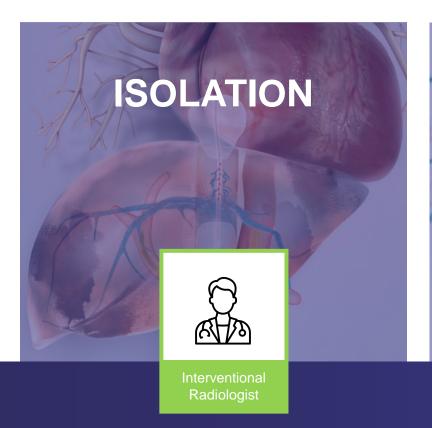
Instructions are given to drink plenty of clear liquids prior to bed the night before the procedure.

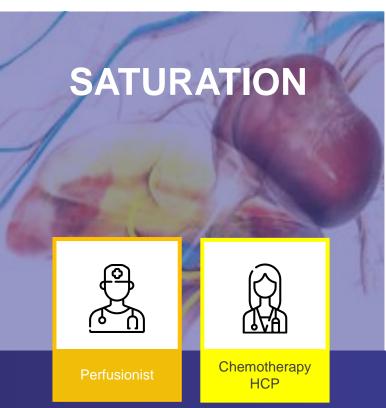
Dehydration increases the risk of unstable blood pressure and difficulty maintaining mean arterial pressures above 65 mmHg.

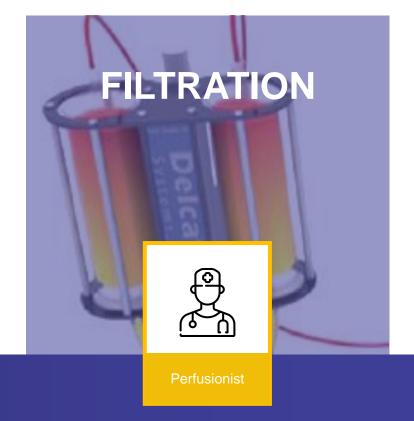
Hydration is initiated at admittance to the hospital on the day of the procedure.

Excessive hydration pre-procedure or intra-procedure can increase post procedure complications. In some cases, excessive hydration may prolong mechanical ventilation due to head and neck edema.

PHP Procedure Team Roles and Responsibilities







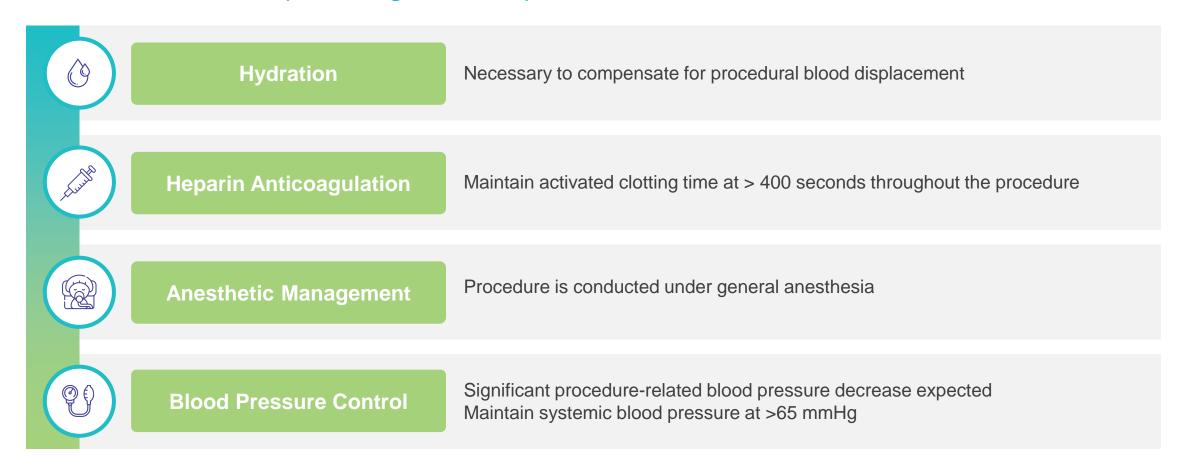
Catheterization Placements

Drug Infusion Administration

Extracorporeal Circuit Filtration Management

Medications that are Administered During the Procedure

All medications and supportive measures must be determined and administered in accordance with each institution's policies, guidelines, procedures.

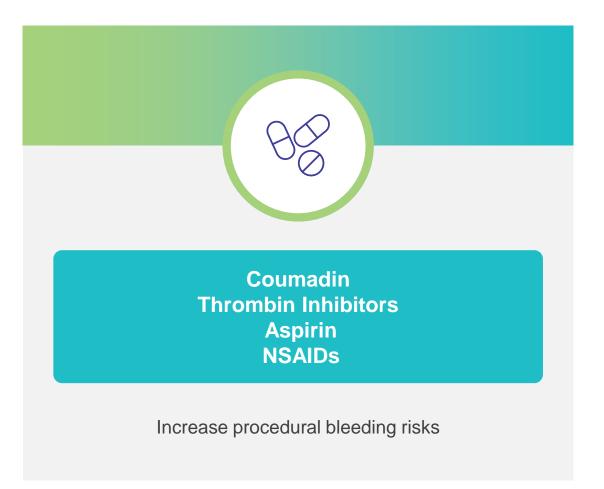


Medications That Should Be Withheld

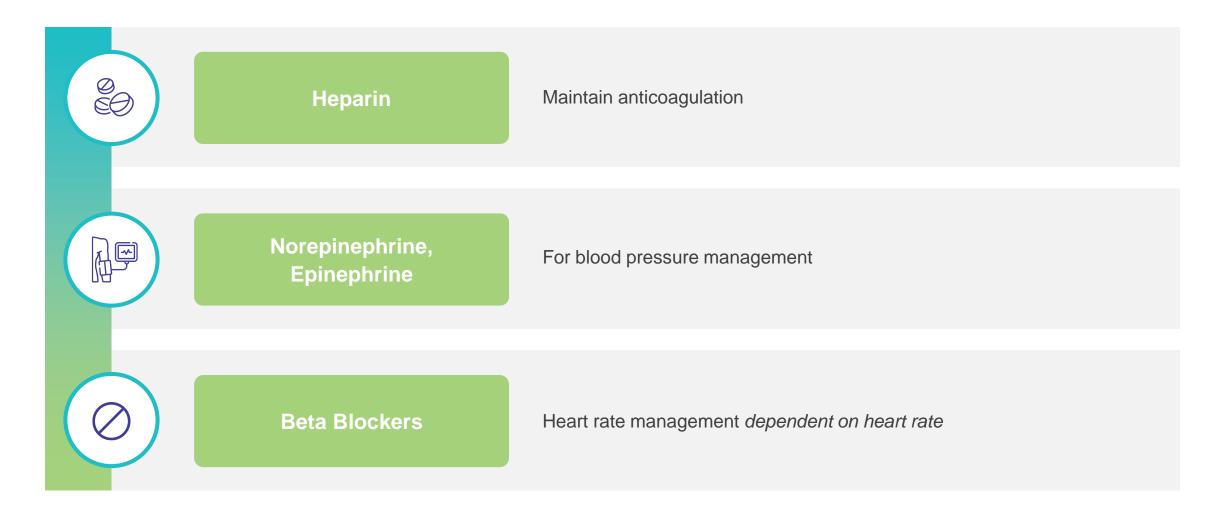


ACE inhibitors
Calcium channel blockers
Alpha 1 adrenergic blockers

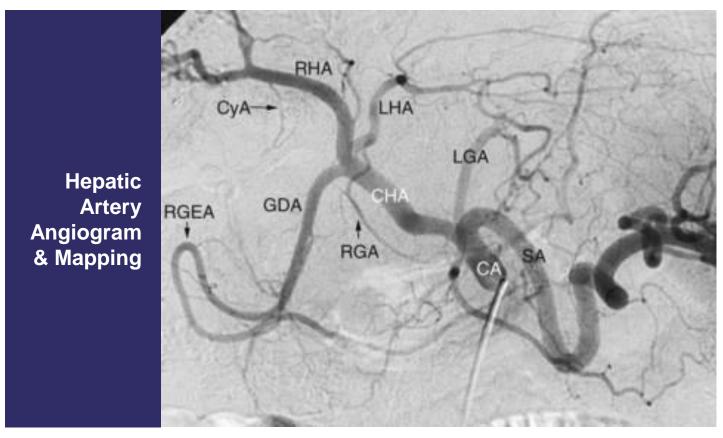
Increase vasopressor resistance



Medications Delivered During the Procedure



Treatment Day Angiogram



Conventional celiac artery anatomy. CA, celiac axis; LGA, left gastric artery; SA, splenic artery; CHA, common hepatic artery; GDA, gastroduodenal artery; RGA, right gastric artery; RHA, right hepatic artery; LHA, left hepatic artery; CyA, cystic artery; RGEA, right gastroepiploic artery.



The arterial supply to the liver must be completely examined the day of the procedure if not previously done one week prior.

Determinations

Possible impact of chemo infusion Embolization needs

PHP Procedure Team Tasks and Key Communications





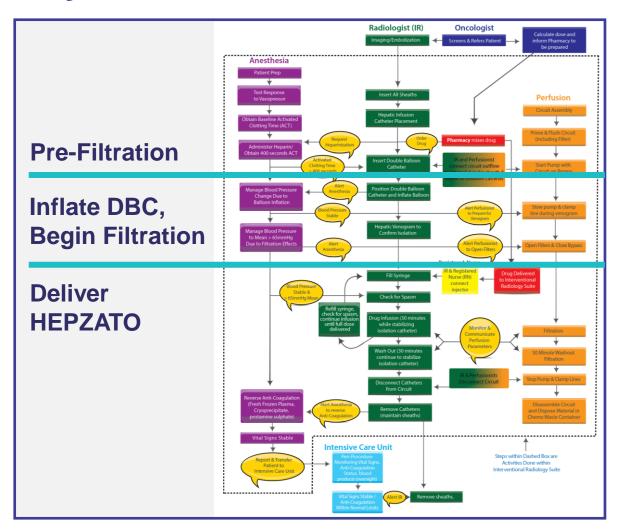


Each PHP Procedure Team member has separate workstreams during the procedure

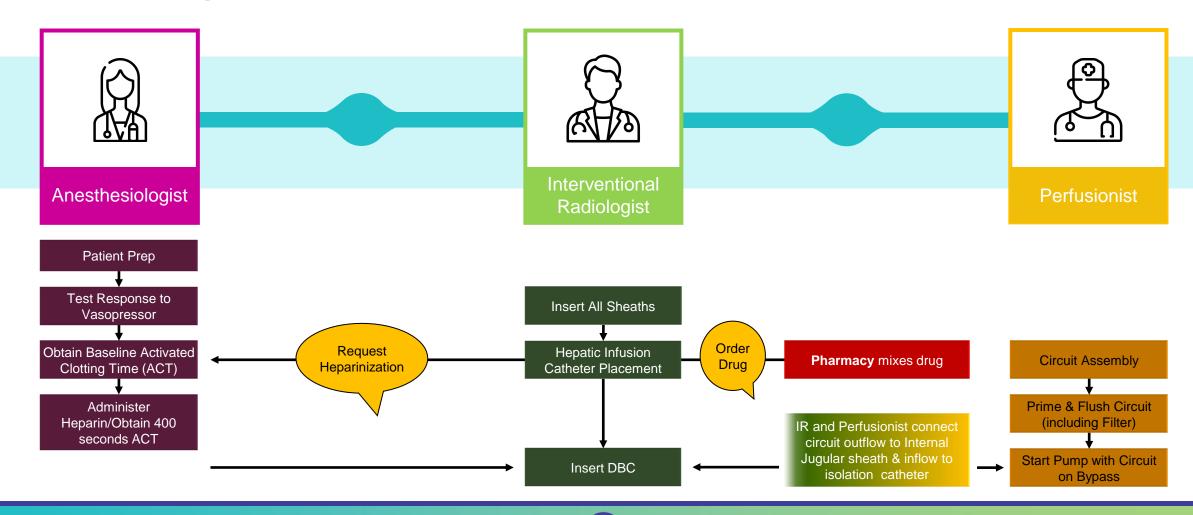
There are critical points during the procedure where key information between the 3 PHP Procedure Team members needs to be shared

Natural pause points in the procedure, where all 3 PHP Procedure Team members must be aligned

the moment before Double Balloon Catheter (DBC) is expanded when extracorporeal filtration begins before drug delivery begins

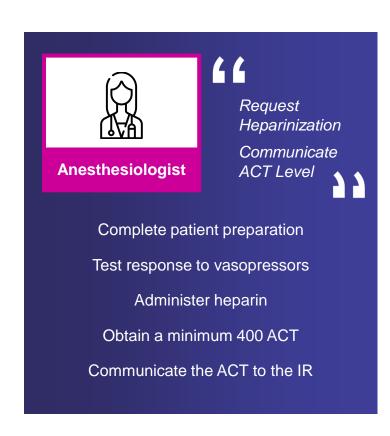


Prepare to Expand Double Balloon Catheter (DBC) and Bring the Filters "on-line"

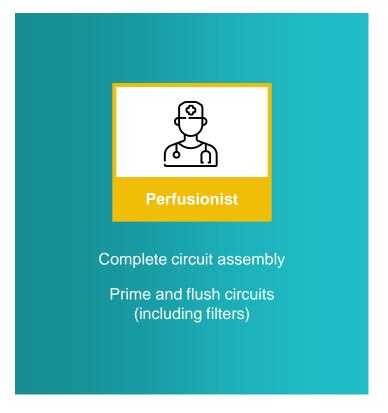


PHP Procedure Team tasks prior to Double Balloon Catheter (DBC) inflation and bringing filters on-line

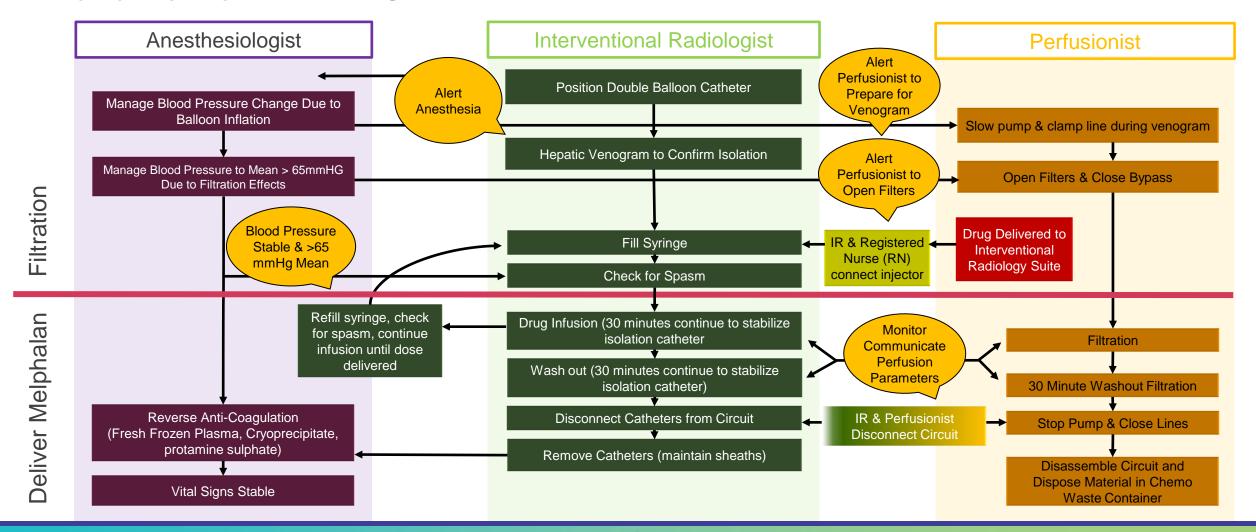
Just prior to bringing the filter "on-line" the 3 PHP Procedure Team members should:



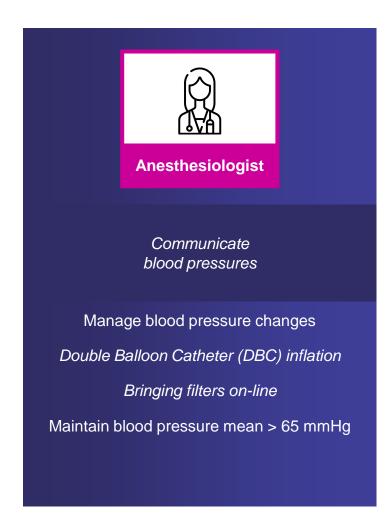


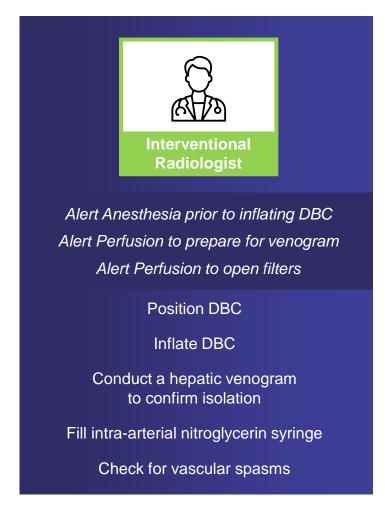


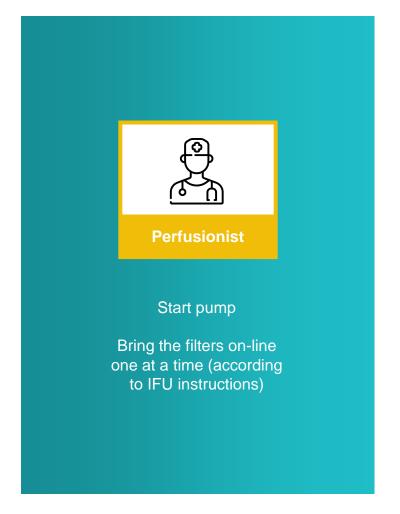
Inflate Double Balloon Catheter (DBC), Bring Filters On-Line, and Deliver HEPZATO



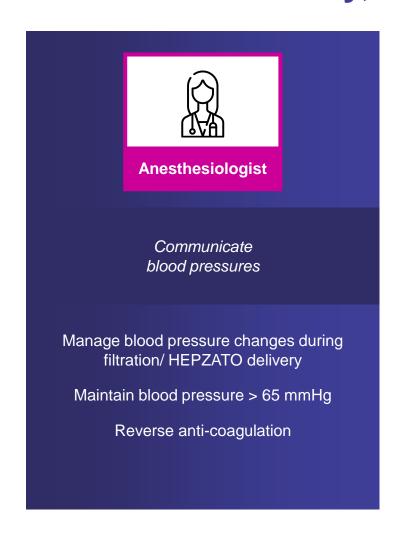
Bringing Filters on-line, Filtration, prior to HEPZATO Delivery

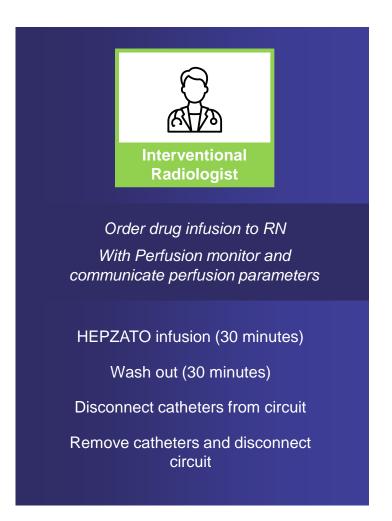


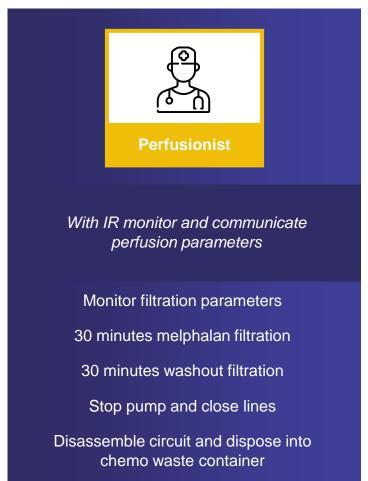




HEPZATO Delivery, Washout and End of PHP Procedure







Perfusionist Procedural Awareness



Pump Flow Rate



The flow rate of the pump should be maintained between 0.4 – 0.8 L/min



Pre pump pressure (suction side) should not be more negative than −250 mmHg, as lower pressures indicate possible catheter collapse or kink.



Pre-cartridge pressures (pre-filter) should not exceed 200 mmHg, as higher pressures indicate increasing filter resistance potentially due to thrombus or a kinked return line. Check filters to assure free flow and return line for kinks.

IR Procedural Flow - Hepatic Artery Spasm

Once the blood pressure is stabilized the IR **injects contrast agent** to check for hepatic artery spasm.

IF SPASM IS OBSERVED:

POSSIBLE CAUSE	0	vasopressor therapy & low mean arterial pressure
EFFECT	0	reflux of melphalan into proximal non-embolized gastrointestinal branches or incomplete delivery of HEPZATO
DURING PROCEDURE	0	perform Angiography every 5 minutes
SPASM RELIEF	•	100 mcg/injection of intra-arterial nitroglycerin in hepatic artery

If there is a persistent intractable spasm, the procedure should be stopped.

Infusion, Filtration & Double Balloon Catheter (DBC) Management

Once the IR approves the start of HEPZATO infusion a 100 mL bolus is administered at 25 mL/minute flow rate using drug injector (estimated 4 min infusion time).

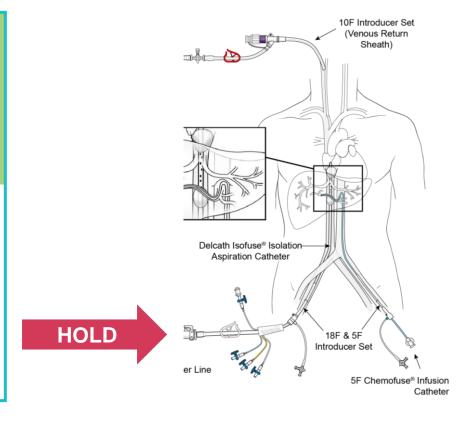
The IR maintains a gentle hold of the DBC position during the entire infusion period.

HEPZATO is refilled into the drug injector and the IR performs a spasm check.



First 30 minutes: HEPZATO Infusion

Last 30 minutes: Washout period



The DBC position checked fluoroscopically every 4 to 5 minutes during drug administration and filtration to ensure continued hepatic venous isolation

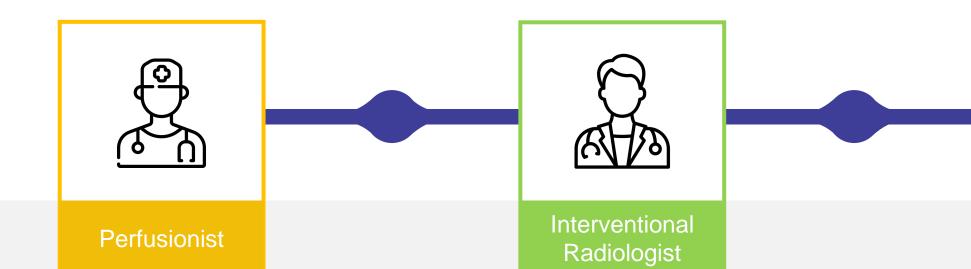
Post PHP Procedure

Blood Return

To return blood to the patient the Interventional Radiologist instructs the Perfusionist to open clamps for saline flush and this pushes the blood through the filter and back into the patient's internal jugular vein.

The effectiveness of the return of blood to the patient is dependent on the central venous pressure and the amount of fluid given.

End of PHP Procedure: Roles and Responsibilities



Stops all Extracorporeal Filtration Circuit (EFC) flow

Opens bypass line

Assists with blood return

Appropriately disposes all components

Collapses Double
Balloon Catheter (DBC)

Disconnects DBC

Disconnects internal jugular introducer from the EFC



Vasopressor weaned/discontinued

Normalizes coagulation

Administers post procedural medications





MODULE 7 Catheterization, Isolation & Saturation

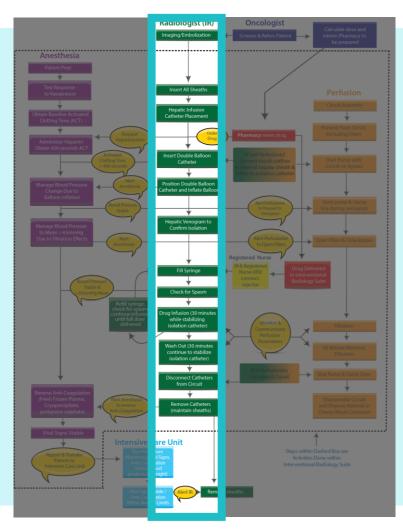
Module 7 Objectives

Module 7 describes the responsibilities of the Interventional Radiologist (IR) on the day of the procedure. This includes catheterization, hepatic isolation, and contrast administration during drug infusion.

- Properly insert and place introducers and catheters.
- Understand procedural techniques that affect filter efficacy and systemic melphalan exposure.
- Confirm hepatic artery anatomy, determining whole liver vs lobar infusion approach and optimal hepatic artery catheter location.
- Assess appropriate Double Balloon Catheter (DBC) position and stabilization.
- Understand process to administer HEPZATO and how to split HEPZATO dose for sequential lobar approach.

Interventional Radiologist Procedural Tasks





This module
describes the steps
the Interventional
Radiologist takes
during the procedure.



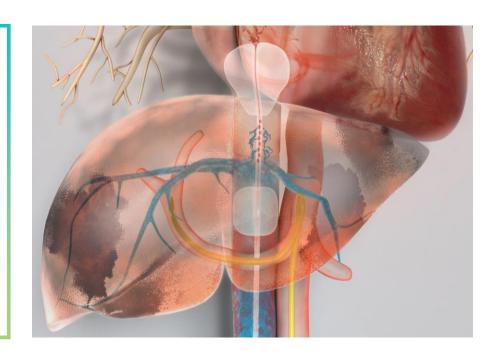
Imaging/Embolization Insert All Sheaths Insert Double Balloon Catheter **Hepatic Infusion Catheter Placement** Position Double Balloon Catheter Hepatic Venogram to Confirm Isolation Fill Syringe **Check for Spasm** Drug Infusion (30 minutes continue to stabilize isolation catheter) Wash out (30 minutes continue to stabilize isolation catheter) Disconnect Catheters from Circuit Remove Catheters (maintain sheaths)





The Interventional Radiologist is the lead physician during the PHP Procedure with HEPZATO KIT and has the following responsibilities:

- Arterial Mapping
- Introducer Placement
- Catheter Positioning
 - The liver blood supply needs to be assessed in order to formulate a strategy for catheter placement.
 - This helps to ensure adequate drug infusion to the entire liver.
 - Dependent on vascular anatomy, this may require repositioning of the catheter during the procedure
- Spasm Management
- Contrast administration during drug infusion



PHP Procedure Eligibility Evaluation



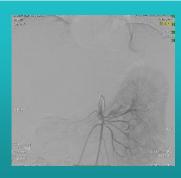
In order to determine eligibility for each patient, the IR evaluates the following:

Prior surgeries that could potentially affect normal hepatic biliary and vascular anatomy

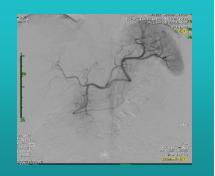
Reimplanted anatomy; such as the common bile duct which can increase risk of biliary tree infection

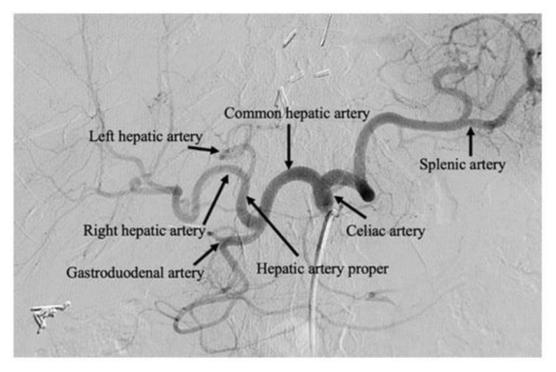
The arterial anatomy is assessed during the mapping process and can consist of a superior mesenteric artery angiogram and a celiac angiogram

Superior Mesenteric Artery Angiogram



Celiac Angiogram





Angiographic evaluation decreases the risk of infusion failure and reflux of melphalan

IR Procedural Awareness



Before the PHP procedure the IR should confirm the following:



- KIT Type Decision
 (balloon spacing 50mm or 62mm)
- Confirm KIT is in the IR Suite
- HEPZATO ordered from the pharmacy
- The patient's most recent lab results
- The treatment start time
- The PHP Procedure team is present



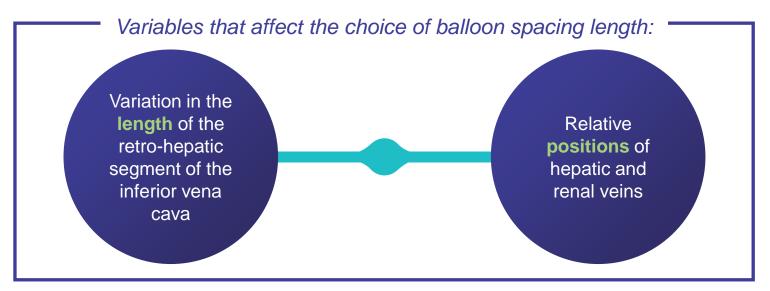
During the PHP procedure the IR must:

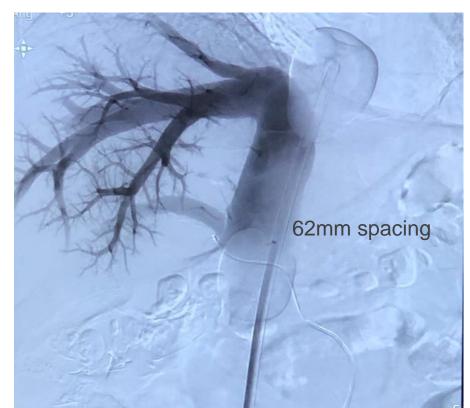
- Lead a PHP Procedure team brief
- Ensure that the Anesthesia team is prepared, and that required medications are at hand
- Confirm that hemofiltration circuit is set up by the Perfusionist
 - Confirm HEPZATO arrival time

Selection of Balloon Spacing



The HEPZATO KIT comes available with two different lengths of balloon spacing: 50mm and 62mm



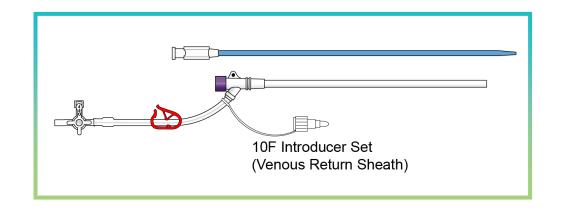


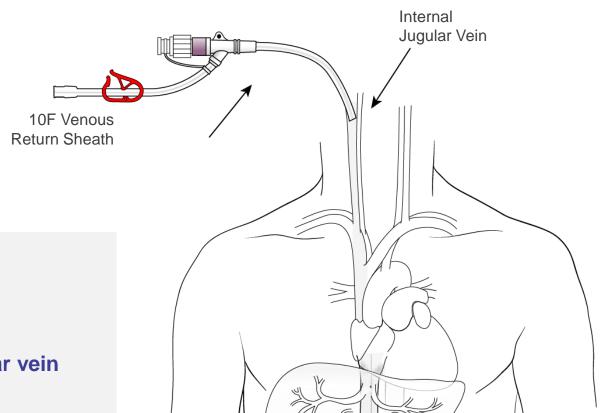
IMPORTANT:

IR needs to decide which KIT (50mm or 62mm spacing) will be used for the PHP procedure before the actual PHP procedure

Interventional Radiologist

Insert 10F Introducer Set







Flush sheath with sterile heparinized saline



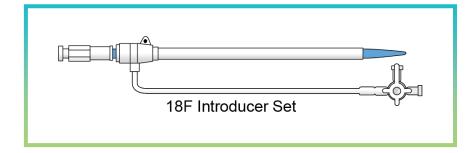
Insert venous return sheath into internal jugular vein



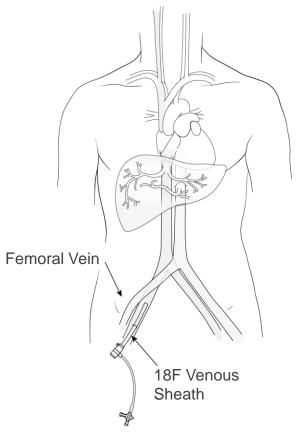
Interventional Radiologist

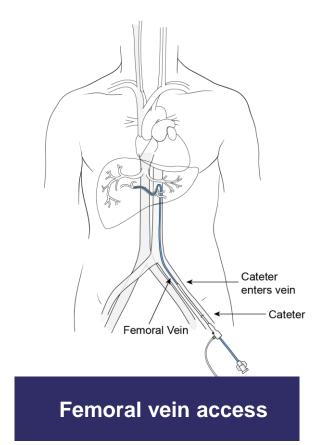
Insert 18F Venous Sheath





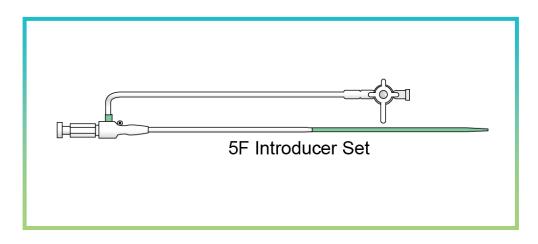
Perform serial dilation with dilators
Insert 18F femoral sheath

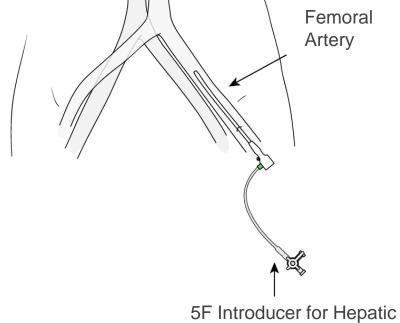




Interventional Radiologist

Insert 5F Introducer Set





Arterial Infusion Catheter



Interventional Radiologist:

"Can heparin be administered?"



Flush 5F sheath



Place 5F sheath into the femoral artery





Hepatic Vascular Mapping

Map
Hepatic
Artery
Anatomy

The Interventional Radiologist needs to create an infusion plan based on hepatic artery mapping.

The infusion plan ensures that all hepatic arteries are adequately perfused.



Embolize As Needed

A selective infusion strategy may be used when the catheter is positioned distal to non-hepatic arterial branches.

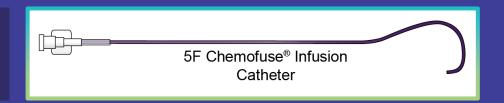


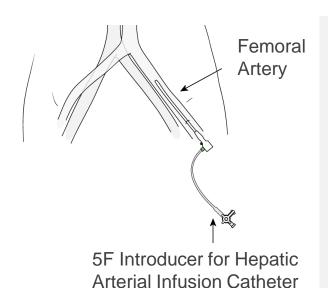




Insert Hepatic Artery Infusion Catheter

Place Hepatic Artery Catheter





The 5F infusion catheter is introduced through the 5F introducer sheath into the femoral artery.

The infusion catheter and microcatheter is placed in the desired position for HEPZATO infusion.



Interventional

"The catheter has been placed."

Radiologist:

Interventional Radiologist

Catheter Location

Based on evaluation of hepatic artery, two different infusion strategies can be utilized: Lobar Infusion or Whole Liver Infusion.

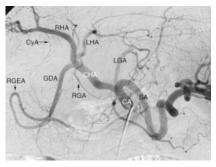
If performing a lobar infusion approach:

- The catheter is positioned to isolate a lobe
- The treatment is administered sequentially
- It requires HEPZATO dose splitting
- Dosing to each lobe is based on relative liver volume and is traditionally a 60% (right lobe) to 40% (left lobe) split.
- The catheter must be repositioned during HEPZATO infusion
- It is suggested to begin with the most difficult catheterization point in order to minimize catheter repositioning time

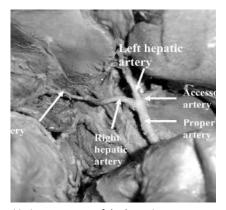
If performing a whole liver infusion approach:

- Performing a whole liver infusion approach is dependent on arterial blood flow
- Whole liver infusion may require minimal embolization.

Distal catheter placement to non-hepatic arterial branches can reduce requirements for embolization. Adequate catheter positioning minimizes risk of infusion failure and reflux.



Embolotherapeutic Strategies for Hepatocellular Carcinoma: 2020 Update, Kishore et al, Cancers



Variant anatomy of the hepatic artery in adult Kenyans, European Journal of Anatomy, 2007, Kitungu et al.



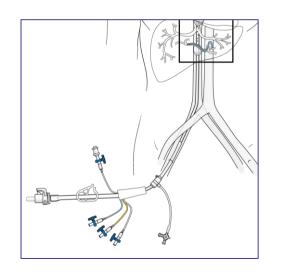


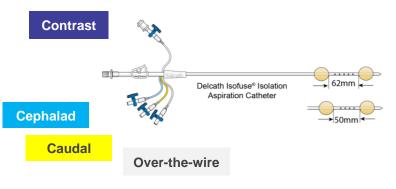
Insert Double Balloon Catheter (DBC)

In preparation of the placement of the DBC:

HEPZATO should be in the room

The ACT must be above 400 seconds





Placement of the DBC:

- The DBC catheter should be flushed with heparinized saline via the contrast port and over-the-wire port and central lumen via the extracorporeal system.
- The DBC catheter is inserted through 18F sheath.



Interventional Radiologist:

"The DBC has been placed"

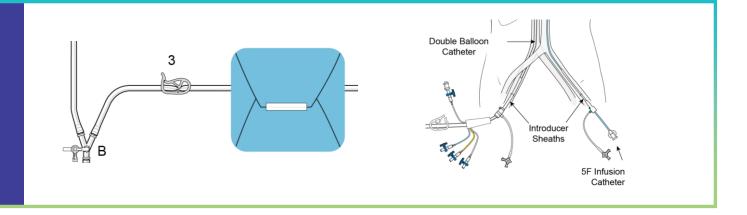




Catheter & Hemofiltration Connection

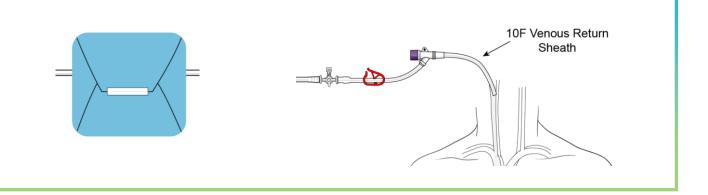
To connect the double balloon catheter to the circuit:

- The sterile wrap needs to be removed from the hemofiltration circuit DBC line.
- A wet connection can be made from the hemofiltration DBC line to the DBC catheter by opening saline clamps 1 and 2 and clamp 3.



To connect the venous return line to the circuit:

- The sterile wrap is removed from the hemofiltration venous return line.
- The hemofiltration circuit is connected to the venous return line to the stopcock of the 10F venous return sheath in the jugular vein.







Establishing Hemofiltration & Isolation

Start ECF Bypass Circuit

When HEPZATO has been delivered to the procedural room and the connection of the hemofiltration circuit to catheters has been made, the perfusionist turns on the pump and then bypass can be started.

BYPASS:

Venous blood aspirated from central lumen

Blood flows through fenestrations in DBC

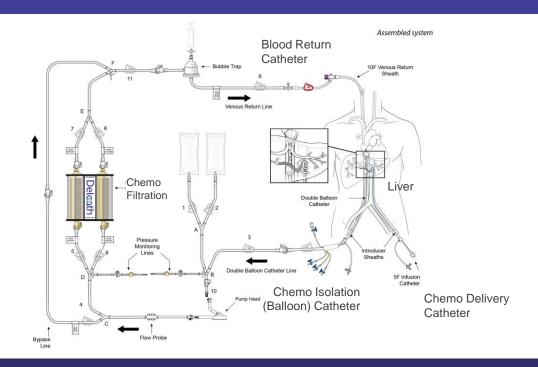
Blood flows through the pump

V

Blood flows through the bypass line

V

Blood returns to the patient through the venous return sheath



During Hemofiltration and Isolation, the Perfusionist continuously monitors the pump pressures



Vasopressor Administration







Prior to occlusion of vena cava and inflation of balloons a vasopressor response test is performed by the Anesthesiologist.



Initial occlusion of the inferior vena cava leads to a significant decrease of systemic blood pressure. It is of critical importance to maintain the blood pressure above 65 mmHg.

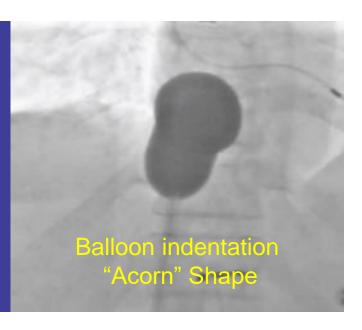


If response to vasopressors is adequate, the double balloon catheter may be inflated.



Balloon Expansion

The balloon is expanded until the lateral edges become effaced with the inferior vena cava wall.



The cephalad balloon must occlude inferior vena cava just *above* the *highest* hepatic vein

The caudal balloon must occlude inferior vena cava just *below* the *lowest* hepatic vein

Never move or reposition the DBC when the balloons are fully expanded.

DO NOT collapse balloons unless HEPZATO administration has been stopped or after completion of infusion, including a 30-minute washout cycle.

The Perfusionist must carefully monitor the circuit flow rate during balloon inflation.

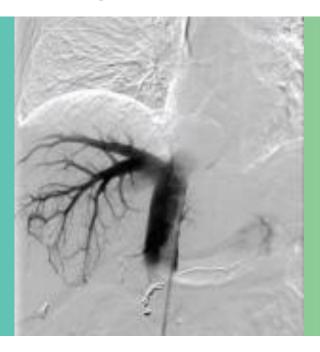


Hepatic Venous Isolation Confirmation & Leak Assessment

lodinated contrast medium is injected through the contrast port to assess isolation of the segment.

To check for balloon leaks:

- 1 The Perfusionist stops the pump.
- The IR checks for leaks by injected contrast agent through the DBC
- If no leak is seen, the Perfusionist starts the pump again.



Hemofiltration Circuit Flow is re-established by:

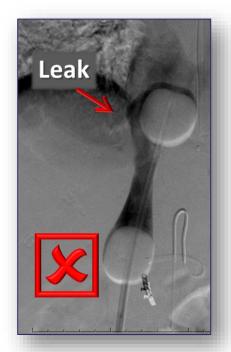
- 1 Restarting the centrifugal pump
- Increasing pump speed to approximately 2200 RPMs or the previously established flow rate

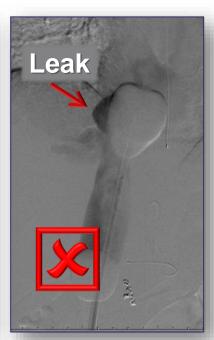
If the **DBC** is not in the proper position, collapse both balloons (caudal balloon first) and then reposition the catheter, while maintaining flow in the hemofiltration circuit





Balloon Occlusion









Balloon Inflation Volumes			
Balloon	Cephalad	Caudal	
Maximum Volume	38 mL	38 mL	

- Under fluoroscopy, the balloons are partially inflated with approximately 15-25 mL of dilute contrast media.
- Inflation of the balloons may give it a rounded appearance.

The balloons must not be over inflated. Overinflation could cause the balloons to burst and result in life-threatening injury.

Op

Opening Filters & Blood Pressure Management



The Process for Turning on the Filters

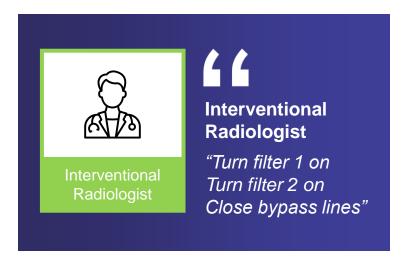
The flow is checked by the IR.

The IR instructs the Perfusionist to turn the filters on; usually one filter at a time.

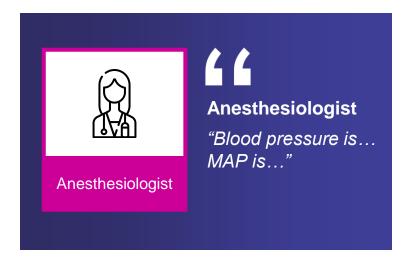
The bypass line is kept open as each filter is opened.

After both filters are filled with blood, the bypass line is closed after 3 minutes of flow.

The blood pressure is monitored for 2 to 5 minutes.









IR Procedural Flow - Hepatic Artery Spasm



Once the blood pressure is stabilized the IR **injects contrast agent** to check for hepatic artery spasm.

IF SPASM IS OBSERVED:

POSSIBLE CAUSE	•	vasopressor therapy & low mean arterial pressure
EFFECT	•	reflux of melphalan into proximal non-embolized gastrointestinal branches or incomplete delivery of HEPZATO
DURING PROCEDURE	•	perform Angiography every 5 minutes
SPASM RELIEF	•	100 mcg/injection of intra-arterial nitroglycerin in hepatic artery

If there is a persistent intractable spasm, the procedure should be stopped.



Infusion, Filtration & Double Balloon Catheter (DBC) Management

Once the IR approves the start of HEPZATO infusion a 100 mL bolus is administered at 25 mL/minute flow rate using drug injector. (estimated 4 min infusion time).

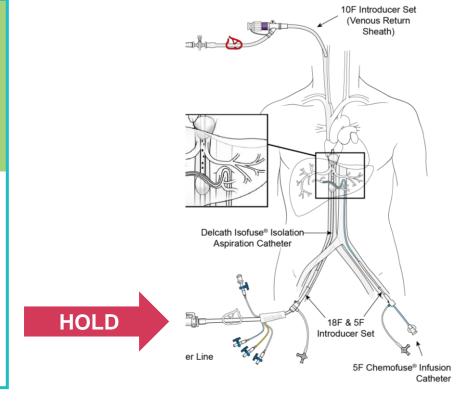
The IR maintains a gentle hold of the DBC position during the entire infusion period.

HEPZATO is refilled into the drug injector and the IR performs a spasm check.



First 30 minutes: HEPZATO Infusion

Last 30 minutes: Washout period



The DBC position checked fluoroscopically every 4 to 5 minutes during drug administration and filtration to ensure continued hepatic venous isolation





Procedure End

The IR may return blood to the patient at the end of the washout period. Blood return may be dependent on central venous pressure and the amount of fluid given.

amount of fluid given.

and this flush pushes the blood through the filter and back into patient's internal jugular vein.

At the end of blood return the Perfusionist properly disposes all the hemofiltration lines, pump, and the filters.

The Anesthesia team starts post procedural medications.

To return blood to the patient the IR instructs

the Perfusionist to open clamps for saline flush



MODULE 8

Anesthesia, Anticoagulation and Hemodynamic Management

Module 8 Objectives

Module 8 describes the responsibilities of the anesthesiologist/anesthetist on the day of the procedure.

This includes medically clearing the patient for the PHP Procedure with HEPZATO KIT, managing preoperative medications, managing anesthesia, blood oxygenation, acid/base balance, core body temperature, hydration, anticoagulation, and blood pressure. Patient to be assessed for adequate cardiopulmonary function.

Managing preoperative medications.

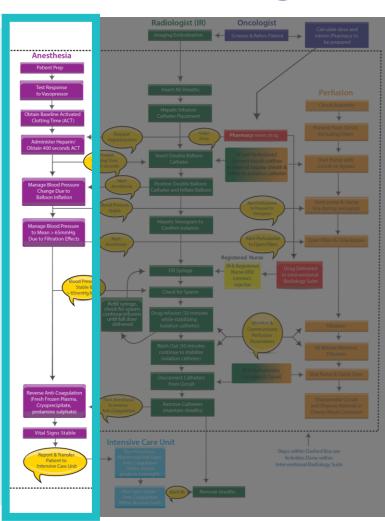
Managing
hydration
optimizing blood
pressure and
EFC flow rate.

Managing anticoagulation.

Managing blood pressure.

Anesthesiologist Procedural Tasks





This module describes the steps that the Anesthesiologist takes during the procedure.



Patient Prep

Test Response to Vasopressor

Obtain Baseline Activated Clotting Time (ACT)

Administer Heparin/Obtain 400 seconds ACT

Manage Blood Pressure Change Due to Balloon Inflation

Manage Blood Pressure to Mean > 65 mmHG Due to Filtration Effects

Reverse Anti-Coagulation (Fresh Frozen Plasma, Cryoprecipitate, protamine sulphate)

Vital Signs Stable

HEPZATO KIT PI



HEPZATO KIT Preoperative Patient Assessment

After referral by the Medical Oncologist, a preoperative assessment of the patient is performed by the Anesthesiologist, the Cardiologist, and the Pulmonologist.

Each HCP must adjudicate pre-operative clearance indicating adequate cardiac and pulmonary function.

A patient may not be eligible for HEPZATO KIT treatment if any of the following are present:

- Active coronary artery disease
- Severe angina
- Recent myocardial infarction
- Any congestive heart failure
- Significant ventricular arrhythmias

- Moderate to severe valvular disease
- Unable to withstand high dose vasopressor therapy
- Advanced COPD
- Unable to withstand mechanical ventilation





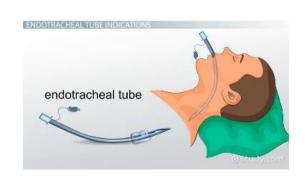


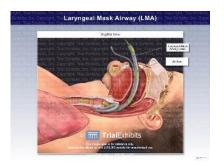
Anesthesia Induction

- A bair hugger can be used to keep the patient warm.
- The Anesthesiology team may use an endotracheal tube or a laryngeal mask to induce anesthesia.

OR







Bair Hugger

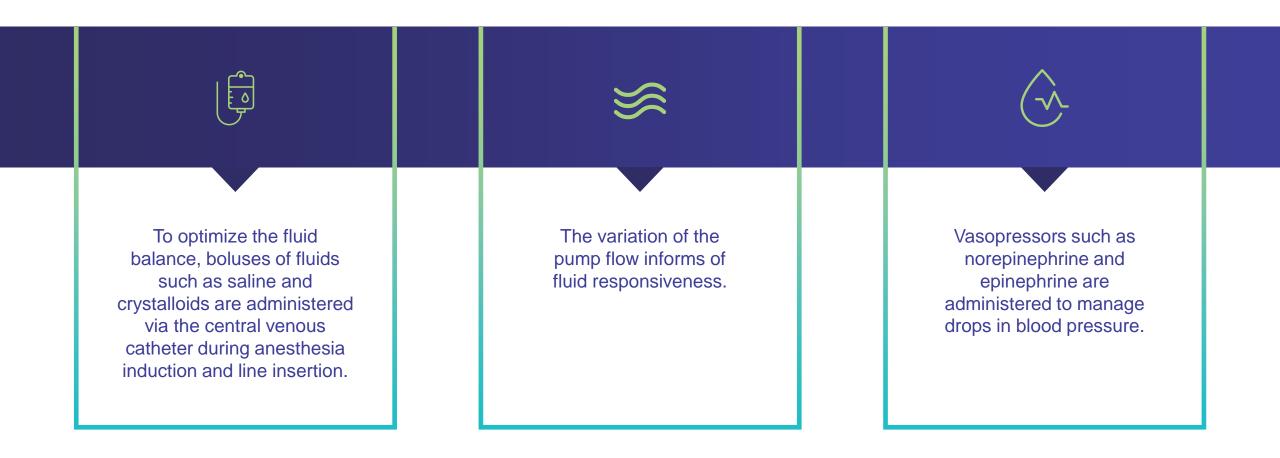
Endotracheal tube

Laryngeal mask





Anesthesia Team





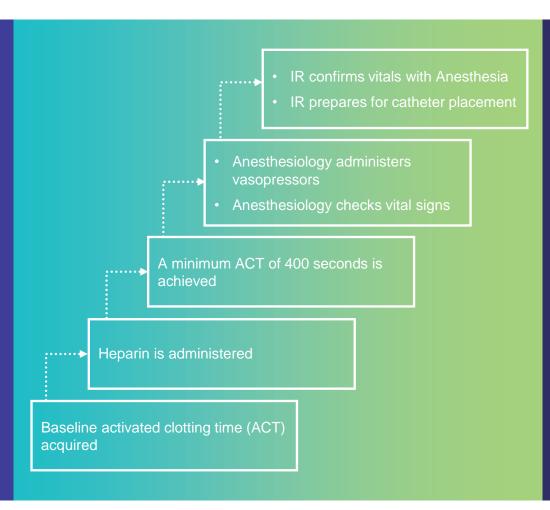




Radiologist:

Interventional

"What is the ACT?"





Anesthesiologist:

"Vasopressors have been administered"

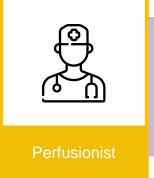
PHP Procedure



Anesthesiologist Role







The Anesthesiologist or the Perfusionist obtains the baseline ACT







- The Anesthesiologist administers Heparin
- The Anesthesiologist administers vasopressors to maintain blood pressure *before* DBC placement by the IR

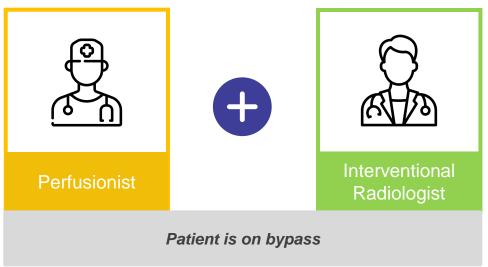
Anticoagulation is required to assure free extracorporeal flow and filtration

PH

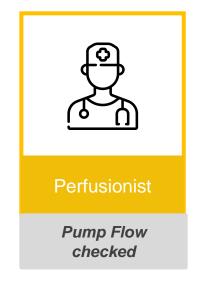
PHP Procedure – Anesthesia Responsibilities



Anesthesia Responsibilities















The **Anesthesiologist:**

Monitors the blood pressure

Administers vasopressors

Stabilizes the mean arterial pressure





Anesthesia Responsibilities



Perfusionist

Filters brought online



Anesthesiologist stabilizes the MAP



Anesthesiologist



Perfusionist

Extracorporeal circulation filtration stopped



Anesthesiologist discontinues vasopressors

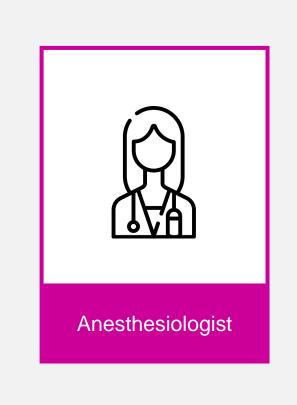


Anesthesiologist



Anesthesiologist

Post Procedure



Discontinues heparin

Initiates recovery

Weans off vasopressors

Removes endotracheal tube or laryngeal mask Administers fluids

Removes arterial catheter and closure device

Extubates the patient

May leave femoral venous sheath and jugular sheath in position

Optional | Give fresh frozen plasma (FFP) and thrombocytes



MODULE 9

Post-Procedure Management & Discharge

Module 9 Objectives

Module 9 describes the responsibilities of the intensivist to recover the patient after the procedure. This includes reversing anticoagulation, removing intravascular introducers (sheaths), normalizing hematologic abnormalities (e.g., anemia and thrombocytopenia), and normalizing procedural abnormalities (e.g., metabolic acidosis and hypothermia).



Mechanisms reversing anticoagulation and safe removal of vascular sheaths.



Correcting hematologic abnormalities.



Correcting metabolic acidosis and hypothermia

NOTE:

Patients should be assessed for severe periprocedural complications associated with HEPZATO KIT for at least 72 hours after the procedure.

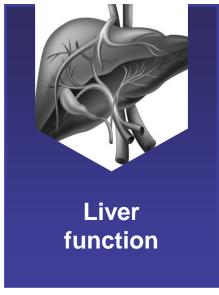
Severe peri-procedural complications should be documented and submitted to the REMS using the Severe Peri-Procedure-related Complications Adverse Events Documentation Form.

Follow up may occur virtually or via telephone for patients discharged prior to 72 hours.

Post Procedural Recommendations

Prior to discharge after the PHP Procedure with HEPZATO, it is recommended that the following procedure-related events are corrected:











Corrections of post-procedure events should follow institutional guidelines and clinical judgment.

End of Procedure

Anticoagulation Interventions & Hematologic Abnormalities

To reverse anticoagulation and attend to procedural induced coagulopathies

Protamine is titrated to normalize the Activated Clotting Time (ACT).

Clotting factors may be replenished by administration of cryoprecipitate and/or Fresh Frozen Plasma.

To correct hematologic abnormalities

Platelets should be replaced according to institutional guidelines.

Ensure that the platelet count is greater than 50,000/mm³ and patient coagulation status is normalized before safely removing sheaths.

Blood product replacement should be considered if patient has significant blood loss. It is suggested to follow institutional guidelines for administration of packed red blood cells, follow anemia hospital protocols, and transfuse accordingly.

Patient Discharge & Follow Up



Discharge

Prior to discharge, physiologic abnormalities should be reviewed and reversed.



Medical Oncologist Responsibilities

Follow up with the patient after hospital discharge.

Clear the patient for subsequent treatment.

Evaluate need for dose adjustment or treatment discontinuation.

Administer prophylactic treatment to manage hematologic toxicities.

Monitor patient on an outpatient basis after hospital discharge for HEPZATO related toxicities