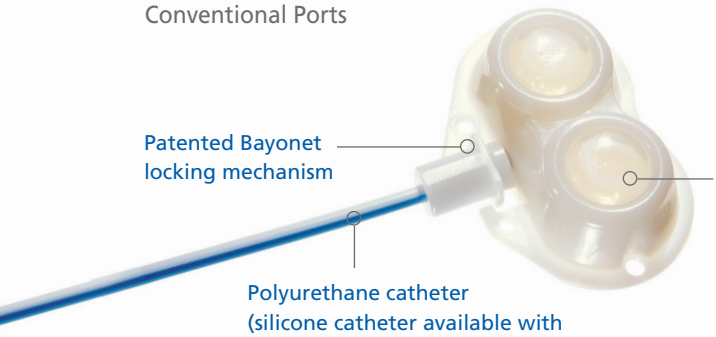
LIFEPORT

CONVENTIONAL PORTS

angiodynamics

LifePort

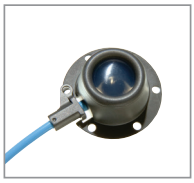
Conventional Ports



Compatibility
Plastic: MRI safe
Titanium: MRI conditional–3 Tesla (available with single port options only)

LIFEPORT CONVENTIONAL PORT COMPLETE SIZE OFFERINGS

LifePort* implantable port body options include plastic and titanium, as well as single, dual and low-profile models. Refer to the complete product list below for more information.



Single Titanium

Single Titanium Port	Introducer Size (F)	Catheter Size (F)	UPN
Detached silicone catheter	10	9.6	H787LPS50130
Detached polyurethane catheter	9	8.4	H787LPS55130
Single Plastic Port	**Introducer Size (F)**	**Catheter Size (F)**	**UPN**
Attached silicone catheter	10	9.6	H787LPS70150
Detached silicone catheter	10	9.6	H787LPS70130
Detached polyurethane catheter	9	8.4	H787LPS75130
Dual Plastic Port	**Introducer Size (F)**	**Catheter Size (F)**	**UPN**
Detached polyurethane catheter	10	9.6	H787LPS75230
Single Low-Profile Titanium Port	**Introducer Size (F)**	**Catheter Size (F)**	**UPN**
Detached silicone catheter	10	8.4/9.6	H787LPS50570
Detached polyurethane catheter	6	5.7	H787LPS55550
Dual Low-Profile Plastic Port	**Introducer Size (F)**	**Catheter Size (F)**	**UPN**
Detached polyurethane catheter	7	6.6	H787LPS72550

PORT SYSTEM INCLUDES: LifePort port system, Catheter, Locking mechanisms (detached models), Introducer needle: 18 Ga, Vein pick, PeelPro* PTFE introducer, Guidewire: 0.035" x 50 cm, Blunt needle (detached models), Tunneler, 10 mL syringes, Infusion Set: 20 Ga x 1"†, Low-Profile Infusion Set: 22 Ga x 0.75"†, Non-coring needle: 22 Ga.

†dual models contain 2 infusion sets, low-profile models contain 22 Ga x 0.75" infusion sets.

IMPORTANT RISK INFORMATION

INDICATIONS FOR USE: AngioDynamics implantable access port systems are intended to facilitate frequent blood sampling or the delivery of medications, nutritions, blood products, and imaging solutions.

CAUTION: Federal (USA) law restricts this device to sale by or on the order of a physician.

CONTRAINDICATIONS: AngioDynamics port systems should not be implanted in the presence of known or suspected infections, septicemia, or peritonitis, in patients who have exhibited prior intolerance to the materials of construction, or patients whose body size or tissue is insufficient to accommodate the size of the port or catheter.

WARNINGS AND PRECAUTIONS: The device is sterile and intended for single patient use. Sterile unless the package is opened or damaged. Do not re-sterilize. Use of AngioDynamics anti-coring (19 to 22 gauge Huber point) needles in all procedures is recommended. Observe all instructions for use. Failure to do so may result in patient complications or device damage.

POTENTIAL COMPLICATIONS: Use of port systems involve potential risks normally associated with the insertion or use of any implanted device or indwelling catheter including but not limited to: Infection; pneumothorax; catheter malposition, migration or fragmentation; catheter pinch-off or rejection; hemorrhage; hematoma; clot formation, thrombophlebitis or thromboembolism; vessel trauma, including puncture, laceration, and erosion of vessel and skin; cardiac arrhythmia, puncture and tamponade; endocarditis; thoracic duct injury; peritonitis; fibrin sheath; and drug extravasation (leakage). Occlusion may result from clot formation inside the lumen of the catheter, precipitate formation inside the port from incompatible drugs, or from catheter tip placement against a vein wall or valve.

Indications, contraindications, warnings and instructions for use can be found in the instructions for use supplied with each device. Observe all instructions prior to use. Failure to do so may result in patient complications.

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