

remedē[®] System

Product Reference Guide

	ITEM	MANUFACTURER	PART/ORDER NUMBER	DESCRIPTION	RECOMMENDED INVENTORY ON HAND
Manufacturer Specific	PG	Respicardia (ZOLL)	1001, 1100, 1600	remedē PG	2
	Lead	Respicardia (ZOLL)	4065, 4165, 4665	respistim LQS, 65cm	2
	Lead	Respicardia (ZOLL)	3102, 3652	respistim R Lead – 24mm dia., normal length	2
	Lead	Respicardia (ZOLL)	3103, 3653	respistim R Lead – 20mm dia., extended length	Reps Carry
	Catheter	Respicardia (ZOLL)	7120-S	respiguide 120°	2
	Catheter	Merit Medical	57538CS-WOR	Impress 5F CS Catheter, 75cm	2
	Catheter	Merit Medical	57538CSV-WOR	Impress 5F CS VERT Catheter, 75cm	2
	Catheter	Merit Medical	1628-043	Impress 5F AZYGOS Right, 75cm	2
	Accessory	Pressure Products	SS-SA-09	9F Sealing Adapter	2
	Accessory	Respicardia (ZOLL)	1007	Test Adapter	Reps Carry
	Accessory	Medtronic	6232ADJ	Adjustable Slitter	2
Physician Preference	Guidewire	Physician preference		0.014" Floppy	3
	Guidewire	Physician preference		0.014" Stiff	2
	Guidewire	Physician preference		0.018"	2
	Guidewire	Physician preference		0.035" angled floppy	2
	Guidewire	Physician preference		0.035" access J-wire	2
	Catheter	Physician preference		9Fr Multipurpose Right Sided Coronary Sinus Guide Catheter	2
	Lead	Physician preference		IS-1 Bipolar CS (LV) Lead, 4Fr dia. preferred	1
	Accessory	Physician preference		Y-connector and hemostasis valve	2
	Accessory	Physician preference		IS-1 standard Port plug	4
	Accessory	Physician preference		8Fr x 13cm hemostatic introducer sheath, splittable	2
	Accessory	Physician preference		Disposable pacing cables	2
	Accessory	Physician preference		Programming Wand Sterile Sleeve	2
	Accessory	Physician preference		Guidewire Torque Device	2

Indication for use: The remedē System is an implantable phrenic nerve stimulator indicated for the treatment of moderate to severe Central Sleep Apnea (CSA) in adult patients. **Contraindications:** The remedē System is contraindicated for use in patients with an active infection or patients known to require magnetic resonance imaging (MRI). See the Instructions for Use for complete information regarding the procedure, indications for use, contraindications, warnings, precautions, and potential adverse events.

Rx Only.

The remedē[®] System, remedē[®] EL System, and remedē[®] EL-X System have received FDA approval. The remedē[®] System model 1001 has received CE Mark approval.

ZOLL MEDICAL CORPORATION

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