remedē® System
Inventory Reference Guide

	ITEM	MANUFACTURER	PART/ORDER NUMBER	DESCRIPTION	MINIMUM INVENTORY QUANTITIES		
					1 CASE	2 CASE	3 CASE
Manufacturer Specific	PG	ZOLL Respicardia	1001, 1100, 1600	**rem**edē PG	2	3	4
	Lead	ZOLL Respicardia	4065, 4165, 4665	respi**stim** LQS, 65cm	2	3	4
	Lead	ZOLL Respicardia	3102, 3652	respi**stim** R Lead – 24mm dia., normal length	2	3	4
	Lead	ZOLL Respicardia	3103, 3653	respi**stim** R Lead – 20mm dia., extended length	1	1	1
	Catheter	ZOLL Respicardia	7120-S	respi**guide** 120°	2	3	4
	Catheter	Merit Medical	57538CS-WOR	Impress 5F CS Catheter, 75cm	2	3	4
	Catheter	Merit Medical	57538CSV-WOR	Impress 5F CS VERT Catheter, 75cm	2	3	4
	Accessory	Pressure Products	SS-SA-09	9F Sealing Adapter	2	3	4
	Accessory	ZOLL Respicardia	1007	Test Adapter (Reps Carry)	2	3	4
	Accessory	Medtronic	6232ADJ	Adjustable Slitter	2	3	4
Physician Preference	Guidewire	Physician preference		0.014" Floppy	3	6	9
	Guidewire	Physician preference		0.014" Stiff	2	3	4
	Guidewire	Physician preference		0.018"	2	3	4
	Guidewire	Physician preference		0.035" angled floppy	2	3	4
	Guidewire	Physician preference		0.035" access J-wire	2	3	4
	Accessory	Physician preference		Y-connector and hemostasis valve	2	3	4
	Accessory	Physician preference		IS-1 standard Port plug (For models 1001, 1100)	3	5	7
	Accessory	Physician preference		8Fr x 13cm hemostatic introducer sheath, splitable	2	3	4
	Accessory	Physician preference		Disposable pacing cables	2	3	4
	Accessory	Physician preference		Programming Wand Sterile Sleeve	2	3	4
	Accessory	Physician preference		Guidewire Torque Device	2	3	4

Indication for use: The **rem**edē System is an implantable phrenic nerve stimulator indicated for the treatment of moderate to severe Central Sleep Apnea (CSA) in adult patients. **Contraindications:** The **rem**edē 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 **rem**edē® System, **rem**edē® EL System, and **rem**edē® EL-X System have received FDA approval. The **rem**edē® System model 1001 has received CE Mark approval.

ZOLL MEDICAL CORPORATION

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