

What's needed for a new referral:

- Demographics
- Detailed Written Order Signed/Dated by physician
- Most recent Face-To-Face note discussing the item the patient is needing as well as the diagnosis



Documentation Checklist Positive Airway Pressure (PAP) Devices

Policy References: <u>Local Coverage Determination Positive Airway Pressure (PAP) Devices (L33718) and Policy Article (A52467)</u>

Documentation References: Standard Documentation Requirements Policy Article (PA) A55426

		lier must be able to provide all of these items on request: ard Written Order (SWO)				
	Refill Requirements					
	Beneficiary Authorization					
	Proof of Delivery (POD)					
	Continued Need					
	Continued Use					
	Medical records from treating practitioner as noted below					
Me	dical F	lecords should contain:				
Init	ial cove	erage (First 3 months)				
Positive Airway Pressure Device - E0601						
		Face to Face (F2F) prior to the sleep test to assess the beneficiary for obstructive sleep apnea (OSA); and				
		Medicare-covered sleep test that meets either:				
		□ Apnea-hypopnea index (AHI) or Respiratory Disturbance Index (RDI) ≥ 15 events per hour with a minimum of 30 events; or				
•		\square AHI or RDI \geq 5 and \leq 14 events per hour with minimum 10 events and documentation of:				
		☐ Excessive daytime sleepiness, impaired cognition, mood disorders, or insomnia; or				
		☐ Hypertension, ischemic heart disease, or history of stroke, and				
		Documentation the beneficiary and/or their caregiver has received instruction from the supplier of the PAP device in the proper use and care of the equipment.				
Do	cumen	tation for Beneficiaries Who Fail the Initial 12 Week Trial				
		F2F re-evaluation to determine the etiology of the failure to respond to PAP therapy; and				

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	Repeat sleep test in a facility-based setting (Type 1 study).			
Bi-level Res	spiratory Assist Device (RAD) without Back-up Rate (E0470)			
	Beneficiary meets all the criteria listed above for a positive airway pressure device (E0601); and			
	An E0601 PAP device has been tried and proven ineffective based on a therapeutic trial conducted in either a facility or home setting.			
	Documented failure to meet therapeutic goals using an E0601 during the titration portion of a facility-based study or during home use despite optimal therapy			
	A new initial F2F if E0601 has been used for more than 3 months and the beneficiary switched to E0470 (a new sleep test is not required)			
PAP – Cont	inued Coverage (Beyond the First 3 Months of Therapy)			
	Documentation the beneficiary is benefiting from PAP therapy as demonstrated by:			
	F2F re-evaluation by the treating physician between the 31st and 91st day after initiating therapy documenting that symptoms of OSA are improved; and			
	Objective evidence of adherence to use of the PAP device reviewed by treating physician.			
	☐ Adherence is defined as use of the PAP device ≥ 4 hours per night on 70% of nights during a consecutive 30 day period anytime during the first 3 months of initial use			
Beneficiarie	es Entering Medicare			
☐ Sleep test – Documentation the beneficiary had a sleep test, prior to FFS Medicare enrollment, meets Medicare AHI/RDI coverage criteria in effect at the time the beneficiary seeks replaceme PAP device and/or accessories; and				
	Clinical Evaluation – Following enrollment in FFS Medicare, the beneficiary must have a F2F which documents:			
	□ Diagnosis of OSA; and			
	☐ The beneficiary continues to use the PAP device.			
Replaceme	nent (E0601, E0470)			
	Replacement following the 5 year reasonable useful life (RUL) requires a F2F that documents the beneficiary continues to use and benefit from the PAP device			
Non-Heated	d or Heated Humidifier (E0561, E0562)			
	Beneficiary meets PAP coverage criteria; and			
	Standard written order includes the type of humidification			



DETAILED PHYSICIANS ORDER RESPIRATORY EQUIPMENT

Patient: Da	te of Order:	HT: W	/T:
Address:	DOB:	Length of Need:	99
	Diagnosis Codes:		•
Equipment Needed:			
E0570 – Nebulizer with compression	on	E0601- CPAP settings:	
Nebulizer Admin Sets		E0470- BiPAP settings:	
A7003 – Admin sets (2/month)	.	E0471 BiPAP ASV settings:	
A7015 – Aerosol Mask (1/month)		E0562 Heated Humidifier	
E0463- Volume Control Ventilator I		CPAP/BiPAP/Respiratory assist :	Supplies:
E0464 – Volume Control Ventilator	Non-Invasive Interface	A7030 – Full Face Mask (1/3r	
E0482 – Cough Stimulating Device		A7031 – Face mask interface	•
E0439 – Stationary Liquid Oxygen S	System	A7032 - Nasal cushions (2/1r	no)
E0463 – Volume Control Ventilator	•	A7033 - Nasal pillows (2/1mg	o) ·
Invasive Interface		A7034 – Nasal application de	vice (1/3mo)
E0464 – Volume Control Ventilator	•	A7035 – Headgear (1/6mo)	
Non-invasive Interface		A7036 – Chinstrap (1/6mo)	
E0482 – Cough Stimulating Device		A7037 – PAP tubing (1/3mo)	
E0439- Stationary Liquid Oxygen Sy	/stem	A7038- PAP filter disposable	(2/1mo)
E0443 – Oxygen contents-gas		A7039- PAP filter non-dispos	able (1/6mo)
E0434 – Portable Liquid Oxygen sys	stem '	A7044- PAP oral interface (1/	′6mo)
E0431 - Gaseous Oxygen Regulator		A7046- Replacement water c	hamber (1/6mo
E1392 – Portable oxygen concentra	ator	A4604- Tubing with heating e	
E1390 - Oxygen Concentrator	•	·	
Oxygen LPM Continuous/Wi		Overnight Pulse Oximet	ry On:
Via Nasal CannulaOR MaskOR Bl	ed into PAP Device	ROOM AIR	
OR Invasive Vent		Oxygen@	lpm .
		CPAP	
TEST ON CONSERVER		BiPAP	
Titrate patient's oxygen setting to achieve			
at rest and during activities of daily living		Other Please Specify:	
oximetry; and set up on the appropriate co			
device or portable oxygen concentrator.			
Physicians			
Name:	Phy	/sicians	
	Sig	nature:	
Address:			
	Da	te:	
Phone:	NP	l:	_
Fax:			