BlueCross BlueShield Minnesota

DURABLE MEDICAL EQUIPMENT CERTIFICATION

BIRMINGHAM SERVICE CENTER • P.O. Box 10527 • Birmingham, AL 35201-0500 Fax: 1-833-719-1603

Check As Appropriate: 🗌 DME 🗌 OXYGEN 🗌 IPPB 🗌 G	LUCOMETER		BIPAP		RECERTIFICATION	
PATIENT INFORMATION COMPLETE ALL ITEMS	PERTAINI	NG TO THE	PATIENT'S	CONDITION AND EQ	UIPMENT	
1. Patient's Name		2. Date Patient Doctor	t Last Seen by	3. Subscriber Nur	mber	
4. Diagnosis				5. Prognosis	oor	
6. Estimated Number of Months Equipment Needed (Do NOT put "INDEFINITE"; be specific) Date Prescribed 8. Rental Period This Certification Applies To (Certification Length CANNOT Exceed 12 Months) First Day (MM-DD-YYYY) Last Day (MM-DD-YYYY) 9. Supplier's Name, Street Address, City, State, ZIP Code, Phone	b. Room Confined? c. Wheelchair Confined? d. Ambulatory?		□ N □ N □ N □ N ? □ N			
10. Supplier's Provider Number	TT. Hequest		0(3)			
GENERAL EQUIPMENT SEE THE SECTIONS	ON THE B	ACK OF THE	F FORM FO	R OXYGEN AND IPP	B	
12. General Equipment Selected for Patient						
□ a. Alternating P.P. & Pump (Complete #15)		COMPLETE W	HEN INDIGAT	ED IN QUESTION 12		
□ b. Bed, Electric (Complete #13 and #14)		13. Regarding	g Electric Beds	s, is the Patient able to w	ork the controls and	
\square c. Bed, Semi-electric (Complete #13 and #14)		cause the	adjustments?)	🗆 Yes 🗆 No	
\square d. Bed, Standard	-	14. Does the	Patient's cond	ition require frequent ch	anges in body	
				an ordinary bed?	о ,	
□ e Bed, Variable Height <i>(Complete #14)</i>		\Box No \Box Yes; condition is:				
□ f. Cane or Quad Cane			,			
□ g. Walker □ With Wheels						
□ h. Wheelchair □ 1) Standard						
□ 2) Electric		15. Does the	Patient now h	ave, or is the Patient		
□ 3) Detachable Arms		susceptib	le to, decubitu	is ulcers?	🗆 Yes 🗆 No	
□ 4) Leg Rests □ 5) Special; Type:			. a. Has the Patient been trained by a Therapist or Physician to use a powered percussor? □			
 □ i. Commode, Bedside □ j. Lift, Patient □ i. Naturizan Hand hald 				at the Patient's home er manual therapy?	🗆 Yes 🗆 No	
□ k. Nebulizer, Hand-held		17. CPAP/BIP/	AP			
□ I. Nebulizer, Ultrasonic		Date of sl	een studv [.]			
□ m. Percussor (Complete #16)						
\Box n. Rails, Bedside						
□ o. Suction Machine		•	ry disturbance			
\Box p. Sitz Bath						
q. Traction Equipment						
🗆 r. Trapeze Bar		BIPAP	pressures:			
□ s. Other (<i>Describe</i>)		18. If for rece	rtification has	Patient demonstrated c	ompliance	
			e of this equip		□ Yes □ No	

SEE REVERSE SIDE FOR SIGNATURE

			sults of the blood gathe carrier in writin							
19. Report Date	PaO ₂ Level (MM of Hg)	Oximetry Level Where Was Test Done? (MM of Hg) □ Patient's Home □ Doctor's Office □ Nursing Home □ Independent Lab □ Hospital			Check Condition of Patient During Oximetry Level Test During Activities, Such as Exerci At Rest			s Patient on Room or Oxygen at Time llood Gas Study? oom Air xygen		
	20. a. Type Oxygen Unit Prescribed: Dortable Stationary Concentrator b. Type Oxygen Unit Prescribed: Liquid Gaseous									
21. How many hours per day is the Patient on Oxygen? a. Non-portable O_2 : hours b. Portable O_2 : hours										
 □ For exercise therapy outside the home:										
23. The following treatments were tried WITHOUT SUCCESS for this Patient PRIOR TO OXYGEN THERAPY:										
					TREA	TMENT DATES:	BEGIN (MM-DD-YYYY)	ENDED (MM-DD-YYYY)		
	□ YES □ NO Bronchodilators:									
	ES INO Medications: MEDICATION NAME			DOSAGE						
□ YES □ NO Phy	vsical Therapy:	a. Percusso								
		🗆 b. Breathing	g Exercises							
□YES □NO Oth										
GENERAL EQUIPMENT CERTIFICATION LENGTH CANNOT EXCEED SIX MONTHS										
24. Current results of any pulmonary function studies are: Forced vital capacity before and after aerosol bronchodilators: 25. What is the IPPB frequency of use?						ncy of use?				
Before	After		redicted V.C.	Date	of Studies	-				
26. IPPB used to <i>(Check all that apply):</i>										
🗆 a. Deliver	aerosolized med	ications			ounteract pulmo		n or edema			
				 f. Decrease the work of breathing g. Regulate inspiratory and expiratory flow patterns 						
\Box d. Correct or prevent atelectasis \Box h. Other (<i>Explain</i>):										
27. Can the Patient successfully use a hand-held nebulizer or a nebulizer with a compressor? \Box YES \Box NO <i>(Explain)</i>										
GLUCOMETER										
28. Is this Patient an insulin-dependent diabetic? $\hfill T YES \hfill NO$				29. What is the average daily dose of insulin? Units						
30. What type of insulin is being used? □ Regular □ NPH 3 □ Other (Describe):					31. What is the number of daily insulin injections?					
32. Does the Patient have widely fluctuating blood sugars before meal time? 33. Does the Patient have frequent episodes of insulin reactions? 33. Does the Patient have frequent episodes of insulin reactions? 33. Does the Patient have frequent episodes of insulin reactions?							n reactions?			
34. a. Is it necessary for the Patient to make frequent checks of his or her blood glucose level? \Box YES \NO b. Is the Patient's Vision impaired enough to require a special glucose monitoring system at home? \Southeas YES \NO c. Is this Patient capable of being trained to use a home blood glucose monitor? \YES \NO										
PHYSICIAN'S INFORMATION, CERTIFICATION OR RECERTIFICATION NOTICE: This form must be completed, signed and dated by the prescribing physician to accurately adjudicate the DME Claim. Any misrepresentation or falsification of information herein may constitute fraud and be subject to legal action.										
34. a. Physician's				-						
	·									
					e Telephone Nu					
35. I certify that I am actively treating this patient, the equipment prescribed is part of my present course of treatment and is "reasonable and necessary," and is not prescribed as convenience equipment, plus all items completed on this form are accurate.										
Attending Physician's Handwritten Signature (STAMPED signature is NOT Acceptable) Date										