

Oxygen & Oxygen Equipment Beneficiaries Meeting Group I Criteria

DOCUMENTATION CHECKLIST

REQUIRED DOCUMENTATION IN SUPPLIER'S FILE

All Claims for Oxygen: Initial Certification

Documentation of Dispensing Order (preliminary written or verbal order) that contains:

Description of the item	Date of the order
Name of the beneficiary	Physician signature (for written order) or supplier signature (for verbal order)
Name of the physician	

NOTE: If the claim includes HCPCS code E0424, E0431, E0433, E0434, E0439, E0441, E0442, E0443, or E0444, a detailed written order must be obtained prior to delivery. This home oxygen equipment cannot be delivered based on a dispensing order. A dispensing order for other equipment related to home oxygen therapy is only required if the items are dispensed prior to obtaining the detailed written order.

Detailed Written Order That Contains:

Beneficiary's name
The treating physician's name
The treating physician's NPI (if claim includes HCPCS code E0424, E0431, E0433, E0434, E0439, E0441, E0442, E0443 or E0444)
The treating physician's signature
The date the treating physician signed the order (personally entered by the physician)
The date of the order
The item(s) to be dispensed – Must include all separately billed accessories/supplies and specify quantity to provide and replacement frequency
The means of oxygen delivery (cannula, mask, etc.)
The flow rate and frequency of use

Physician's signature on the written order meets **CMS Signature Requirements**

<http://www.cgsmedicare.com/jc/pubs/news/2010/0410/cope12069.html>

Certificate of Medical Necessity for Home Oxygen (The CMN may act as a substitute for a written order if it is sufficiently detailed)

Proof of Delivery

Beneficiary's name	Serial number
Quantity delivered	Delivery date
Detailed description of item(s)	Signature of person accepting delivery
Manufacturer	Relationship to beneficiary

Medical records supporting that the patient meets the basic coverage criteria specified in the coverage and payment rules section of the Oxygen and Oxygen Equipment LCD

The treating physician has determined that the patient has a severe lung disease or hypoxia-related symptoms that might be expected to improve with oxygen therapy, **AND**
The patient has had a blood gas study that meets one of the following criteria, **AND:**

At rest (awake but sitting or lying down), the arterial PO₂ is at or below 55 mm Hg
or the arterial oxygen saturation is at or below 88%.

While awake, the patient's arterial PO₂ is \geq 56 mm Hg or the arterial oxygen saturation is \geq 89% but, for at least 5 minutes during sleep, the arterial PO₂ falls to \leq 55 mm Hg or the arterial oxygen saturation to \leq 88%. During sleep, there is a decrease in the arterial PO₂ of more than 10mm Hg or a decrease in the arterial oxygen saturation of more than 5% from baseline saturation for at least 5 minutes **and** the decrease in PO₂ or O₂ saturation is associated with symptoms or signs reasonably attributable to hypoxemia.

NOTE: The value reported on the CMN must be the lowest value (not related to artifact) during the 5 minute qualifying period. See the LCD (<http://www.cgsmedicare.com/jc/coverage/LCDinfo.html>) for complete details on the rules regarding home sleep oximetry studies.

Baseline saturation = Mean saturation level during the duration of the test.

For beneficiaries with OSA, a qualifying oxygen saturation test for the purpose of determining Medicare home oxygen reimbursement may only occur during a titration polysomnographic study. Please refer to the Positive Airway Pressure Devices and Oxygen Local Coverage Determinations (LCD) ([http://www.cms.gov/medicare-coverage-database/indexes/lcd-list.aspx?Cntrctr=140&name=CGS%20Administrators,%20LLC%20\(18003,%20DME%20MAC\)&DocType=Active&ContrVer=2&CntrctrSelected=140*2&LcNtrctr=140*2&bc=AgACAAIAAAAAA%3d%3d&#ResultsAnchor](http://www.cms.gov/medicare-coverage-database/indexes/lcd-list.aspx?Cntrctr=140&name=CGS%20Administrators,%20LLC%20(18003,%20DME%20MAC)&DocType=Active&ContrVer=2&CntrctrSelected=140*2&LcNtrctr=140*2&bc=AgACAAIAAAAAA%3d%3d&#ResultsAnchor)) and the online article, "Frequently Asked Questions: Oxygen Use in Beneficiaries with Obstructive Sleep Apnea" (<http://www.cgsmedicare.com/jc/pubs/news/2013/1113/cope23913.html>) for additional information.

At rest, the patient's arterial PO₂ is \geq 56 mm Hg or the arterial oxygen saturation is \geq 89% on room air **but**, during exercise, the arterial PO₂ falls to \leq 55 mm Hg or the arterial oxygen saturation is \leq 88% **and**, oxygen administration improves the hypoxemia **and**, medical record includes **all** of the following:

- Blood gas study performed at rest without oxygen;
- Blood gas study performed during exercise without oxygen; and
- Blood gas study performed during exercise with oxygen applied that demonstrates improvement of the hypoxemia.

NOTE: All three qualifying blood gas study reading should be taken during a single testing session. The blood gas reading obtained during exercise, while breathing room air, is the number that should be recorded on the CMN. However, all three readings must be recorded in the medical record and available to the DME MAC or other Medicare contractors upon request.

The qualifying blood gas study was performed by a physician or by a qualified provider or supplier of laboratory services (blood gas studies performed by a supplier are not acceptable), **AND**

The qualifying blood gas study was obtained under one of the following conditions:

Performed during an inpatient hospital stay, no earlier than 2 days prior to the hospital discharge date, and was the last test obtained prior to discharge; or
Was not performed during an inpatient hospital stay and was performed while the patient was in a chronic stable state, not during a period of acute illness or an exacerbation of their underlying disease, **AND**

The qualifying blood gas study was the most recent study obtained prior to the Initial Date indicated in Section A of the CMN and this study was obtained within 30 days prior to the Initial Date, **AND**

The patient was seen and evaluated by the treating physician within 30 days prior to the date of initial certification, **AND**

Alternative treatment measures have been tried or considered and deemed clinically ineffective.

Physician's signature on the written order meets **CMS Signature Requirements**
<http://www.cgsmedicare.com/jc/pubs/news/2010/0410/cope12069.html>

Recertification (Required 12 months after Initial Certification)

Recertification CMN

Medical records documenting that the patient was seen and re-evaluated by the treating physician within 90 days prior to the date of the recertification.

- * If the physician visit is not obtained within the 90-day window but the beneficiary continues to use oxygen and the visit is obtained at a later date, coverage would resume beginning with the date of that visit. The date of the visit is the recertification date that must be entered on the Recertification CMN.

Please refer to the **LCD** ([http://www.cms.gov/medicare-coverage-database/indexes/lcd-list.aspx?Cntrctr=140&name=CGS%20Administrators.%20LLC%20\(18003.%20DME%20MA%20C\)&DocType=Active&ContrVer=2&CntrctrSelected=140*2&LCntrctr=140*2&bc=AgACAA-IAAAAAA%3d%3d&#ResultsAnchor](http://www.cms.gov/medicare-coverage-database/indexes/lcd-list.aspx?Cntrctr=140&name=CGS%20Administrators.%20LLC%20(18003.%20DME%20MA%20C)&DocType=Active&ContrVer=2&CntrctrSelected=140*2&LCntrctr=140*2&bc=AgACAA-IAAAAAA%3d%3d&#ResultsAnchor)) for complete details regarding when an initial, recertification, or revised CMS is required.

Continued medical need for the equipment/accessories/supplies is verified by either:

- A refill order from the treating physician dated within 12 months of the date of service under review; or
- A change in prescription dated within 12 months of the date of service under review; or
- A properly completed CMN with an appropriate length of need specified; or
- A medical record, dated within 12 months of the date of service under review that shows usage of the item.

Portable Oxygen Systems

Medical records that support:

- The patient is mobile within the home; and
- The qualifying blood gas study was performed at rest (awake) or during exercise

Liter flow greater than 4 LPM

- A copy of a blood gas study showing blood gas levels in the Group I or Group II range while the patient was receiving oxygen at the rate of 4 LPM

Additional Information References on the Web

- Supplier Documentation Requirements: <http://www.cgsmedicare.com/jc/pubs/pdf/Chpt3.pdf>
- Local Coverage Determinations (LC Ds) and Policy Articles: <http://www.cgsmedicare.com/jc/coverage/LCDinfo.html>
- Oxygen Resources: http://www.cgsmedicare.com/jc/coverage/mr/Oxygen_Resources.html

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