

# CERT

DME MAC OUTREACH & EDUCATION

Task Force for Error-Free Medicare Claims

## Documentation Requirements

June 18, 2024

**noridian**  
Healthcare Solutions

  
**CGS**<sup>®</sup>  
A CELERIAN GROUP COMPANY

# Disclaimer

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The DME MAC CERT Outreach and Education Task Force consists of representatives from each of the DME MACs and is independent from the CMS CERT Team and CERT Contractors, who are responsible for the calculation of the Medicare Fee-for-Service Improper Payment Rate.

The DME MAC CERT Outreach and Education Task Force has produced this material as an informational reference for providers furnishing services in our contract jurisdictions. The CERT Task Force employees, agents, and staff make no representation, warranty, or guarantee that this compilation of Medicare information is error-free and will bear no responsibility or liability for the results or consequences of the use of this material. Although every reasonable effort has been made to assure the accuracy of the information within these pages at the time of publication, the Medicare program is constantly changing, and it is the responsibility of each provider to remain abreast of the Medicare program requirements. Any regulations, policies and/or guidelines cited in this publication are subject to change without further notice. Current Medicare regulations can be found on the Centers for Medicare & Medicaid Services (CMS) website at <http://www.cms.gov>.

# Webinar Moderators and Presenters

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- Jurisdiction A and D – Noridian Healthcare Solutions
  - Shelly Carlson
- Jurisdiction B and C – CGS Administrators
  - Judie Roan

# Agenda

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- Comprehensive Error Rate Testing (CERT) Data
- Documentation Requirements
- Resources
- Questions

# Comprehensive Error Rate Testing (CERT) Data



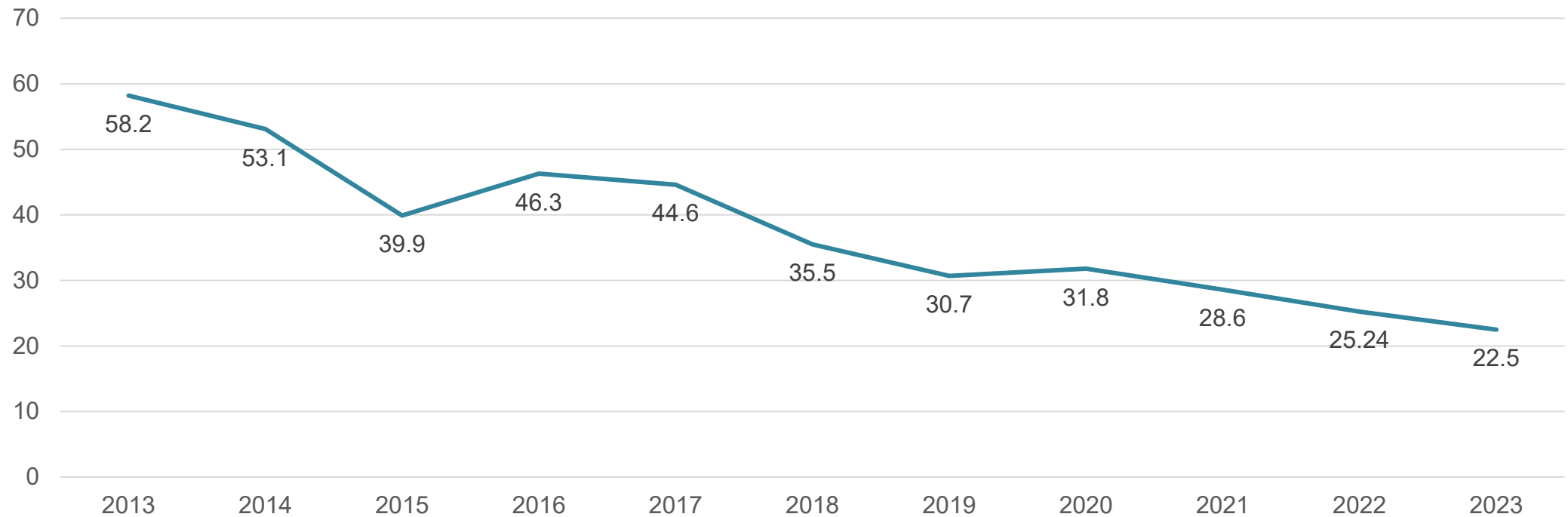
# CERT

2023 Improper Payment Rates and Projected Improper Payment

<https://www.cms.gov/files/document/2023medicarefee-servicesupplementalimproperpaymentdatapdf.pdf>

Service Type	Improper Payment Rate	Projected Improper Payment Amount
Overall	7.4%	\$31.2 B
★ DMEPOS	★ 22.5%	★ \$1.9 B
Part A (excluding Hospital Inpatient Prospective Payment System (IPPS))	7.8%	\$14.2 B
Part A (Hospital IPPS)	3.4%	\$4.1 B
Part B Providers	10.0%	\$11.0 B

# DMEPOS Improper Payment Rate All DME MAC Trend




# Top Root Causes of Improper Payments

Root Cause Description
Documentation to support coverage criteria - missing/inadequate
Documentation to support continued medical need - missing
Order - missing/inadequate
Order not written by provider listed on the claims as ordering/ referring provider
Refill request - missing/inadequate
Units of service (UOS) ordered does not support the units of service (UOS) provided and billed
Proof of delivery - missing/inadequate
Proof of delivery the date of delivery was not supported by the submitted documentation



# CERT Letter

  
CENTERS FOR MEDICARE & MEDICAID SERVICES

PROVIDER/SUPPLIER NAME  
ADDRESS LINE 1  
ADDRESS LINE 2  
CITY, STATE, ZIPCODE

Date: 4/8/2022  
Reference ID: CID #: 00000000  
NPI Provider #: 0000000000  
Phone: 000-000-0000  
Fax: 000-000-0000

**Request Type & Purpose: First Letter**  
**Subject: Additional Documentation Required.**

Dear Medicare Provider/Supplier,

The Centers for Medicare & Medicaid Services (CMS), through the Comprehensive Error Rate Testing (CERT) program, carries out the task of requesting, receiving, and reviewing medical records.<sup>1</sup> The CERT program reviews selected Medicare A, B and DME claims and produces annual improper payment rates. For more information regarding the CERT program, please visit [www.cms.gov/CERT](http://www.cms.gov/CERT).

**Reason for Selection**  
The CMS' CERT program has randomly selected one or more of your Medicare claims for review.

**Action: Medical Records Required**  
Federal law requires that providers/suppliers submit medical record documentation to support claims for Medicare services upon request. Providers/suppliers are required to send supporting medical records to the CERT program. **Providing medical records of Medicare patients to the CERT program does not violate the Health Insurance Portability and Accountability Act (HIPAA).** Patient authorization is not required to respond to this request. Providers/suppliers are responsible for obtaining and providing the documentation as identified on the attached Bar Coded Cover Sheet. The CMS is not authorized to reimburse providers/suppliers for the cost of medical record duplication or mailing. If you use a photocopy service, please ensure that the service does not invoice the CERT program.

**When: 5/23/2022**  
Please provide the requested documentation by 5/23/2022. A response is still required by 5/23/2022 even if you are unable to locate the requested information.

**Consequences**  
If the provider/supplier fails to send the requested documentation or contact CMS by 5/23/2022, the provider's/supplier's Medicare contractor will initiate claims adjustments or overpayment recoupment actions for these undocumented services.

<sup>1</sup>Social Security Act Sections 1833 [42 USC §1395l(e)] and 1815 [42 USC §1395g(a)]; 42 CFR 405.980-986

CENTERS FOR MEDICARE AND MEDICAID SERVICES  
CERT DOCUMENTATION CENTER  
8701 Park Central Drive  
Suite 400-A  
Richmond, VA 23227

**Important Dated Information Enclosed**

**Immediate Response Required**  
**Medicare Record Request**

If no addressee name is shown, forward to Medical Records Department.

# CERT Documentation Requests

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- The CERT Review Contractor requests and receives all medical records
- Use the barcoded coversheet as your documentation coversheet
- Documentation may be submitted:
  - Mail or Fax
  - Electronic Submission of Medical Documentation (esMD)
    - Include a CID number or Claim number and the barcoded cover sheet in your file transmission
    - Information on esMD can be found at <https://www.cms.gov/esMD>
  - CD or Email attachment
    - encrypted per HIPAA security rules
- Check the current status of a claim under CERT review by using the CERT C3HUB Claim Status Search <https://c3hub.certrc.cms.gov/>
  - If CERT shows the review has been completed – refer to DME MAC CERT resources

# CERT Contact Information

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## CERT Contractor Resources and Contacts

- Phone: 1.888.779.7477 or 1.443.663.2699
- E-mail: [certprovider@empower.ai](mailto:certprovider@empower.ai)
- Website: C3HUB (<https://c3hub.certrc.cms.gov/>)

## DME MAC CERT Resource Locations

- JA: <https://med.noridianmedicare.com/web/jadme/cert-reviews/cert>
- JB: <https://www.cgsmedicare.com/jb/claims/cert/index.html>
- JC: <https://www.cgsmedicare.com/jc/claims/cert/index.html>
- JD: <https://med.noridianmedicare.com/web/jddme/cert-reviews/cert>

# Appeal Rights from CERT Audits

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- If the CERT contractor finds errors with the claim in question, the supplier will receive an Overpayment Demand Letter and a revised Medicare Remittance Advice (MRA).
- If the supplier does not agree with the outcome of the CERT review, they should file an appeal to the Redeterminations department of their DME MAC within 120 days of the date on the demand letter or MRA.
  - If a Redetermination is filed to the appropriate DME MAC within 30 days of the letter/MRA, all recoupment activities will cease until the redetermination decision is made.

# Documentation Requirements



# Medical Records

## CERT Errors

Documentation to support coverage criteria – **missing or inadequate**

Documentation to support continued medical need – **missing**



# Documentation Requirements

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- Standard Written Order (SWO) or Written Order Prior to Delivery (WOPD)
- Beneficiary Authorization
- Advance Beneficiary Notice of Noncoverage (ABN) (when applicable)
- Refill Documentation (when applicable)
- Information required for use of specific modifiers
- Clinical documentation to support medical need and continued use of the item
- Proof of Delivery

# Medical Records

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- The medical records include:
  - The treating practitioner's office records
  - Hospital records
  - Nursing home records
  - Home health agency records
  - Records from other healthcare professionals and test reports
- Records from suppliers or healthcare professionals with a financial interest in the claim outcome are not considered sufficient by themselves for determining that an item is reasonable and necessary
- Supplier-produced records, even if signed by the treating practitioner, and attestation letters (e.g., letters of medical necessity) are deemed not to be part of a medical record for Medicare payment purposes



# Documentation in the Beneficiary's Medical Record

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- Should substantiate the medical necessity for the item, quantity ordered, and frequency of use
- Should include (but **is** not limited to):
  - Beneficiary's diagnosis
  - Duration of condition
  - Clinical course
  - Prognosis
  - Nature and extent of functional limitations
  - Other therapeutic interventions and results
  - Past experience with related items, etc.
- All documentation must be maintained in your files for seven years and be available upon request

# CMS Signature Requirements

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- Services provided/ordered/certified must be identifiable by the persons responsible for the care of the beneficiary in accordance with Medicare’s policies:
  - Handwritten
  - Electronic
  - Stamp signatures are typically not acceptable unless the practitioner has physical disability.
- Medicare contractors shall consider the totality of the medical record when reviewing
- CMS Signature Requirements:
  - CMS Program Integrity Manual 100-8, Chapter 3, Section 3.3.2.4:  
<https://www.cms.gov/regulations-and-guidance/guidance/manuals/downloads/pim83c03.pdf>
  - MLN Fact Sheet “Complying with Medicare Signature Requirements”  
<https://www.cms.gov/mln905364-complying-with-medicare-signature-requirements>

# Initial Need Documentation

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- Initial justification for medical need is established at the time items are first ordered
- Medical records demonstrating that the items are reasonable and necessary are created just prior to, or at the time of, the creation of the initial prescription

# Documentation Continued Need/Continued Use Supplies and Rented Items

## Continued Need

TIMELY: Within the preceding  
12 months (unless specified per policy)

Continued Need Documentation
A recent order by the treating practitioner for refills
A recent order/prescription by the treating practitioner for repairs
A recent change in the order/prescription
Timely documentation in the medical record showing usage of items

## Continued Use

TIMELY: Within the preceding  
12 months (unless specified per policy)

Continued Use Documentation
Timely documentation in the medical record showing usage of the item(s), related <b>options</b> /accessories, and supplies
Supplier records documenting requests for refill/replacement of supplies in compliance with the refill documentation requirements (this is deemed to be sufficient to document continued use for the base item as well)
Supplier records documenting beneficiary's confirmation of continued use of a rental item

# Amendments and Corrections to Medical Records

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- All services are expected to be documented in the medical record at the time they are rendered.
- If services provided were not properly documented, documentation may need to be amended, corrected or entered after rendering the service.
- Medical records amended, corrected, or entered after the service should note:
  - The date and author and
  - Change/addenda should be clearly and permanently denoted
- CMS Program Integrity Manual 100-8, Chapter 3, Section 3.3.2.5:  
<https://www.cms.gov/Regulations-and-Guidance/Guidance/Manuals/Downloads/pim83c03pdf.pdf>

# Orders

CERT Errors

Missing or Inadequate

Submitted order not written by provider listed on the claims as ordering/  
referring provider



# Orders/Prescriptions

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- All claims require a written order/prescription from the treating practitioner
  - Written order/prescription = Standard Written Order (SWO)
  - Written Order Prior to Delivery (WOPD) = SWO received prior to delivery
  - Required face-to-face encounter and written order prior to delivery list
    - <https://www.cms.gov/files/document/required-face-face-encounter-and-written-order-prior-delivery-list.pdf>

# Required Face-to-Face and Written Order Prior to Delivery List

- CMS maintains the Required Face-to-Face Encounter and Written Order Prior to Delivery (WOPD) List: <https://www.cms.gov/files/document/required-face-face-encounter-and-written-order-prior-delivery-list.pdf>
  - Forty-six Power Mobility Devices (PMDs) are on the list as dictated by statute: K0800 - K0808, K0813 - K0816, K0820 - K0829, K0835 - K0843, and K0848 - K0864
  - Spinal Orthosis: L0631, L0637, L0648, and L0650
  - Knee Orthosis: L1832, L1833, L1843, and L1851
  - Ankle-Foot/Knee - Ankle Foot Orthosis: L1932, L1940, L1951, L1960, L1970, L2005, and L2036
  - Shoulder/Wrist Hand Orthosis: L3960
  - Osteogenesis Stimulator: E0748
- Items on this list require:
  - Face-to-Face Encounter within six months preceding the order and
  - Written Order Prior to Delivery (WOPD)



# Updates to the Written Order Prior to Delivery (WOPD) and Face-to-Face Encounter (F2F) List

On May 13, 2024, CMS announced updates to the Master List and Required Face-to-Face (F2F) Encounter and Written Order Prior to Delivery (WOPD)

- **Effective August 12, 2024**, CMS selected additional codes that require a F2F Encounter and WOPD:
  - Lumbar-Sacral Orthoses
    - L0635, L0636, L0638, L0639, L0640, L0651
  - Knee Orthoses:
    - L1845, L1852
  - Hospital Beds:
    - E0290, E0301, E0304
  - Osteogenesis Stimulators:
    - E0747, E0760
- Effective August 12, 2024, **L1833 (knee orthosis) no longer requires a WOPD and F2F**

# Written Order Prior to Delivery (WOPD)

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A WOPD is a completed SWO that is communicated to the supplier before delivery of the item

- Items appearing on the Required List require a Face-to-Face (F2F) encounter and WOPD
- The date of the WOPD shall be on or before the date of delivery
  - Must be completed within 6 months after the required F2F encounter
  - For items other than Power Mobility Devices (PMDs), someone other than the treating practitioner may complete certain required elements of the SWO
- F2F encounter and WOPD requirements are statutorily required for PMDs
  - Must be completed within six months before the date on the written order/prescription
  - The WOPD must be completed by the treating practitioner
- MLN Matters Number: SE20007 <https://www.cms.gov/files/document/se20007.pdf>

# Face-to-Face Encounter

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- The treating practitioner must document and communicate to the DMEPOS supplier that they had a face-to-face encounter with the patient within the 6 months before the date on the written order/prescription
  - The 6-month timing requirement doesn't replace other CMS policies
- In-person or telehealth encounter between the treating practitioner and the patient
  - Telehealth encounter must meet the requirements of 42 CFR 410.78 and 42 CFR 414.65
- Supporting documentation includes subjective and objective information associated with diagnosing, treating, or managing a clinical condition for the DMEPOS item ordered
- Suppliers must maintain the written order/prescription, and the supporting documentation provided by the treating practitioner to support payment for the item(s) of DMEPOS and make them available to CMS or its contractors upon request

# SWO/WOPD Elements

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A SWO must include all required elements:

- Beneficiary's name or Medicare Beneficiary Identifier (MBI)
- Order date
- General description of the item
- Quantity to be dispensed, if applicable
- Treating practitioner name or NPI
- Treating practitioner's signature

# SWO/WOPD Description

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- The description can be either a general description (e.g., wheelchair or hospital bed), a HCPCS code, a HCPCS code narrative, or a brand name/model number.
- For equipment - In addition to the description of the base item, the SWO may include all concurrently ordered options, accessories, or additional features that are separately billed or require an upgraded code (list each separately).
- For supplies - In addition to the description of the base item, the order/prescription may include all concurrently ordered supplies that are separately billed (List each separately).

# Entering the Correct Ordering Practitioner on Claims

To help reduce CERT errors:

- When there is more than one practitioner involved in the beneficiary's care:
  - Enter the ordering practitioner's information on the claim
    - Enter the practitioner's name and National Provider Identifier (NPI)
      - > This is the practitioner that created the order

Order	
Name: Jane Doe	Start date: 02/12/24
Equipment: L0650	
Quantity: 1	
Referred by: Dr. John Wilson	
Ordered by: <i>Dr. Jessica Smith</i> NPI: 1234567890	

17. NAME OF REFERRING PROVIDER OR OTHER SOURCE		17a.	
DK	Jessica Smith	71b.	NPI 1234567890

# Refill Request

CERT Errors

Missing or Inadequate



# Refill Documentation Requirements

## Refill Documentation Requirements for dates of service prior to January 1, 2024

Obtained In Person @ Retail Store	Written Request From Beneficiary	Telephone Contact Between Supplier and Beneficiary
Signed delivery slip or copy of itemized sales receipt	Beneficiary name and/or authorized rep ( <i>indicate relationship</i> )	Beneficiary name and the name of person contacted ( <i>if someone other than the beneficiary include this person's relationship to the beneficiary</i> )
Delivery slip/receipt should indicate items were picked up		
	Date of request	Date of contact
	Description of each item requested	Description of each item requested
	Quantity/functional condition of each item still remaining	Quantity/functional condition of each item still remaining
	Contact no sooner than 14 calendar days prior to delivery/shipping	Contact no sooner than 14 calendar days prior to delivery/shipping
	Shipment/delivery occur no sooner than 10 calendar days prior to current supply exhausting	Shipment/delivery occur no sooner than 10 calendar days prior to current supply exhausting



# Refill Requests

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Medicare Program Integrity Manual (PIM) Chapter 5 Section 5.2.6:

<https://www.cms.gov/regulations-and-guidance/guidance/manuals/downloads/pim83c05.pdf>

For refill requests on or after January 1, 2024:

- Suppliers must obtain documentation of beneficiary's affirmative response indicating a need for the refill
  - Removed: Suppliers must document quantity/functional condition of each item remaining
- Suppliers must document the beneficiary has confirmed their need for refill no sooner than 30 calendar days prior to the expected end of the current supply
  - Removed: Contact with the beneficiary or designee regarding refills must take place no sooner than 14 calendar days prior to the delivery/shipping date

# Refill Documentation Requirements: Final Rule CMS1780-F

Refill Documentation Requirements for dates of service **on or after** January 1, 2024

Obtained In Person @ Retail Store	Delivered Refill Communications
Signed delivery slip or copy of itemized sales receipt	Beneficiary name and/or authorized representative ( <i>Suggested: if someone other than the beneficiary include this person's relationship to the beneficiary</i> )
Delivery slip/receipt should indicate items were picked up at store front	
	Date of request
	Description of each item requested
	Documentation of affirmative response indicating a need for the refill
	Contact must occur no sooner than 30 calendar days prior to the expected end of the current supply
	Shipment/delivery occur no sooner than 10 calendar days prior to expected end of current supply

# Proof of Delivery

CERT Errors

Missing or Inadequate

The date of delivery was not supported by the submitted documentation



# Proof of Delivery Methods

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- **Three methods of delivery:**
  - Method 1 – Direct Delivery to the Beneficiary by the Supplier
  - Method 2 – Delivery via Shipping or Delivery Service Directly to a Beneficiary
  - Method 3 – Delivery to Nursing Facility on Behalf of a Beneficiary
- **Regardless of the method of delivery, the contractor must be able to determine that the item(s) delivered are the same item(s) submitted for Medicare reimbursement and that the item(s) were received by a specific beneficiary.**
- Delivery by the supplier
  - Suppliers, their employees, or anyone else having a financial interest in the delivery of the item(s) are prohibited from signing and accepting an item on behalf of a beneficiary (i.e., acting as a designee on behalf of the beneficiary)

# Method 1: Direct to Beneficiary

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Proof of Delivery (POD) must include:

- Beneficiary's name
- Delivery address
- The quantity delivered
- A description of the item(s) being delivered. The description can be either a narrative description (e.g., lightweight wheelchair base), a HCPCS code, the long description of a HCPCS code, or a brand name/model number
- Date delivered
- Beneficiary (or designee) signature

**DATE OF SERVICE = DATE OF DELIVERY**

# Method 2: Shipping Service

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The Proof of Delivery (POD) must include:

- Beneficiary's name
- Delivery address
- Delivery service's package ID number, supplier invoice number or alternative method that links supplier's delivery documents with delivery service's records
- A description of the item(s) being delivered. The description can be either a narrative description, a HCPCS code, the long description of a HCPCS code, or a brand name/model number.
- Quantity delivered
- Date delivered
- Evidence of delivery

**DATE OF SERVICE = SHIPPING DATE,  
LABEL CREATION DATE, OR DATE OF  
DELIVERY**

# Valid Proof of Delivery Example: Method 2 Shipping Service

Shipping Invoice has beneficiary's name, delivery address, description of item(s), quantity shipped

Tracking slip has delivery address, date shipped, and date delivered

**DELIVERY TICKET**  
Date 12/05/2022  
Sales Order

Customer: DOB      Height      Weight      Sex F

Bill to      Deliver to: **Beneficiary's name and delivery address redacted**

Insurance  
Medicare

Comments or Special Instructions  
Tracking number redacted

Delivered On:  
12/05/2022

Delivery Date	Time	CSR	Branch
12/05/2022			

Qty	UOM	Type	Bin	Item
Warehouse				
2	PCS	OTS	Purchase	FreeStyle Libre 14-day Sensor / 71940n R0553

**ups** Proof of Delivery

Dear Customer,

This notice serves as proof of delivery for the shipment listed below.

**Tracking Number**  
Tracking number redacted

**Weight**  
0.40 LBS

**Service**  
UPS Ground

**Shipped / Billed On**  
12/05/2022

**Delivered On**  
12/08/2022 11:44 A.M.

**Received By**

**Delivered To**  
City and State redacted

**Left At**  
Front Door

Thank you for giving us this opportunity to serve you. Details are only available for shipments delivered within the last 120 days print for your records if you require this information after 120 days.

Sincerely,  
UPS

Tracking results provided by UPS: 12/08/2022 1:30 P.M. EST

The tracking number links invoice to tracking slip

# Method 3: Delivery to Skilled Nursing Facility

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Proof of Delivery (POD) must include:

- Documentation demonstrating delivery of the item(s) to the facility by the supplier or delivery entity; and,
- Documentation from the nursing facility demonstrating receipt and/or usage of the item(s) by the beneficiary. The quantities delivered and used by the beneficiary must justify the quantity billed.

**DATE OF SERVICE = SHIPPING DATE OR DATE OF DELIVERY**



# Equipment Retained From a Prior Payer

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A POD is required for all items submitted to Medicare, even those in the beneficiary's possession provided by another insurer prior to Medicare eligibility

- The supplier record must document:
  - A statement, signed and dated by the beneficiary (or beneficiary's designee), that the supplier has examined the item, meets the POD requirements; and
  - A supplier attestation that the item meets Medicare requirements

# Thank You!

