



2025 Coding and Billing Guide



For additional assistance in coding and billing visit reprisereimbursement.com or call our Reimbursement Hotline at 888-249-6793

Disclaimer: This information is for educational/informational purposes only and should not be construed as authoritative. The information presented here is current as of January 2025 and is based upon publicly available source information. Codes and values are subject to frequent change without notice. The entity billing Medicare and/or third party payors is solely responsible for the accuracy of the codes assigned to the services or items in the medical record. When making coding decisions, we encourage you to seek input from the American Medical Association (AMA), relevant medical societies, the Centers for Medicare and Medicaid Services (CMS), your local Medicare Administrative Contractor, and other health plans to which you submit claims. Items and services that are billed to payors must be medically necessary and supported by appropriate documentation. It is important to remember that while a code may exist describing certain procedures and/or technologies, it does not guarantee payment by payors.

Healthcare Common Procedural Coding System (HCPCS), Current Procedural Terminology (CPT) and International Coding Diagnosis and Procedural (ICD-10) coding is universal; however, coverage and payment of medical and surgical services can vary among government (Medicare, Medicaid, TRICARE) and private or Commercial health insurance companies. The procedures described in this guide are widely covered by government and commercial insurers when MiroDry and MiroDerm Wound Matrix is applied in hospitals (both inpatient and outpatient), ambulatory surgery centers (ASCs), and other clinic-based practices. Accurate coding is important to guide how MiroDry and MiroDerm Wound Matrix, and the surgical procedures it is used with, are covered and paid appropriately. Commercial insurer payment methodologies and payment levels will vary from Medicare and other government payers' payment methodologies. It is not possible to capture all insurers' payment methodologies in this guide. Providers should contact their patients' specific insurers directly to obtain information on unique billing, coverage, and payment requirements.

Reprise Biomedical

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Customer Service: 952-377-8238 customerservice@reprisebio.com



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MiroDry is a dry, open, and porous collagen sheet matrix designed to conform to wound beds. It is derived from highly vascularized porcine liver that has been perfusion decellularized, dried, and packaged.

Miro Derm[®]

Biologic Wound Matrix

MiroDerm is a non-crosslinked acellular wound matrix that is derived from the highly vascularized porcine liver. The fenestrations on the product offer a left-to-right stretch which increases the surface area available to contact the wound. It is perfusion decellularized and packaged in an inner sterile pouch with phosphate buffered saline and outer non-sterile pouch.

MiroDry and MiroDerm are sterile medical devices that should be stored in a clean, dry location at room temperature, in the original package. Avoid prolonged exposure to elevated temperatures as it may compromise device functionality. The products expiration dates are indicated as year (4 digits), month (2 digits) and day (2 digits).

Both MiroDry and MiroDerm wound matrix are indicated for the management of wounds, including: partial and full-thickness wounds; pressure ulcers; venous ulcers; chronic vascular ulcers; diabetic ulcers; tunneled, undermined wounds; trauma wounds (abrasion, lacerations, partial thickness burns, skin tears); drainage wounds; and surgical wounds (donor sites/grafts, post-Mohs surgery, post-laser surgery, podiatric, wound dehiscence). See Instructions For Use (IFU) for full prescribing information, including indications, contraindications, precautions, and potential complications.

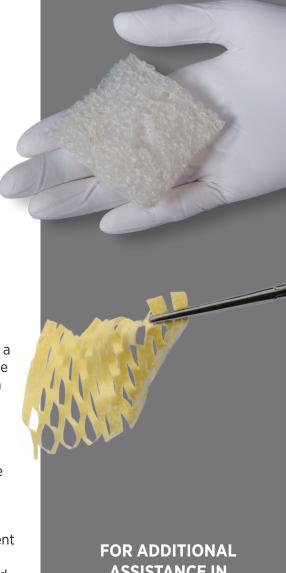
PLACING AN ORDER

Email: customerservice@reprisebio.com

Phone: 952-377-8238 Fax: 952-856-5085

Delivery time: Two business days from receipt of purchase order. Use the following guidelines:

- Orders received by 3 p.m. Central Time will be shipped via FedEx 2-Day Delivery (The customer can elect to have product shipped FedEx Priority Overnight and pay the shipping difference)
- Thursday shipments will be scheduled for delivery on the following Monday
- Friday shipments will be scheduled for delivery on the following Tuesday
- Please call Customer Service with urgent requests.



ASSISTANCE IN CODING AND BILLING



Reimbursement **Hotline Phone & Fax**

P: 888-249-6793 F: 763-317-1977



R³ Reimbursement **Resources Portal**

RepriseReimbursement.com

MiroDry Product Coding

Physicians Office - Use A2031 to document **per square centimeter**. Document MiroDry use in Box 19 of the CMS-1500 Form or cost center description, if payer required.

HCPCS	DESCRIPTION	
A2031	MiroDry Wound Matrix, per square centimeter	
MODIFIERS	DESCRIPTION	
JC	Skin Substitute used as graft	
JW	Drug amount discarded/not administered to any patient	
JZ	No discarded amount: full amount administered	

Important Billing Instructions¹:

- MiroDry is not included on the Medicare Part B Average Sales Price (ASP) File published quarterly by the Centers for Medicare and Medicaid Services (CMS).
- Medicare "A2XXX" HCPCS codes are carrier priced in the non-facility setting or high or low-cost categorized in the facility settings.
- Please reference your specific Medicare Administrative Contractor (MAC) or specific insurer for payment rates and modifier requirements.

Use either JW or JZ Modifiers to detail wastage or lack of wastage²:

Please reference your specific Medicare Administrative Contractor (MAC) or patient-specific insurance carrier for instructions on billing wastage for skin substitutes identified by HCPCS "A" codes.

CMS requires that providers and suppliers report the JW or JZ modifier on Medicare Part B claims for discarded drugs and biologicals. Also, providers and suppliers must document the amount of discarded drugs or biologicals in Medicare beneficiaries' medical records including:

- Date, time, and location of ulcer treated
- Name of skin substitute and how product supplied
- Approximate amount of product unit used
- Approximate amount of product unit discarded
- Reason for the wastage
- Manufacturer's serial/lot/batch or other unit identification number on graft material

The amount discarded should be billed on a separate line with the JW modifier. The unit field should reflect the amount of tissue discarded. The modifier is not required if no discarded portion is being billed to any payer.

 $Please \ refer \ to: \ \underline{https://www.cms.gov/Outreach-and-Education/Medicare-Learning-Network-MLN/MLNMattersArticles/Downloads/MM9603.pdf \ and \ the FAQs \ at \ https://www.cms.gov/Medicare/Medicare-Fee-for-Service-Payment/HospitalOutpatientPPS/Downloads/JWModifier-FAQs.pdf$

Use modifier JZ on billing claims to attest there was no discarded amount from the single-dose vial or single-use package that is normally paid under Part B. Reference (IOM 100-4 Chapter 17, Sections 40-40.1) or page 621 of the CMS CY2023 OPPS/ASC Final Rule: https://public-inspection.federalregister.gov/2022-23918.pdf

^{1.} https://www.cms.gov/medicare/payment/part-b-drugs/asp-pricing-files and www.cms.gov/files/document/b2-2024-public-meeting-agenda-day-2-november-7-2024.pdf

Please refer to: https://www.cms.gov/Outreach-and-Education/Medicare-Learning-Network-MLN/MLNMattersArticles/Downloads/MM9603.pdf and the FAQs at https://www.cms.gov/Outreach-and-Education/Medicare-Learning-Network-MLN/MLNMattersArticles/Downloads/MM9603.pdf and Use modifier JZ on billing claims to attest there was no discarded amount from the single-use package that is normally paid under Part B. Reference (IOM 100-4 Chapter 17, Sections 40-40.1) or page 621 of the CMS CY2023 OPPS/ASC Final Rule: https://public-inspection.federalregister.gov/2022-23918.pdf. No changes were made to this instruction in Federal Register updates for CY2024 or CY2025. Please check with your specific MAC for additional instructions.

MiroDerm Product Coding

Physicians Office - Use Q4175 to document per square centimeter.

HCPCS	DESCRIPTION	
Q4175	MiroDerm Wound Matrix, per square centimeter	
MODIFIERS	DESCRIPTION	
JC	Skin Substitute used as graft	
JW	Drug amount discarded/not administered to any patient	
JZ	No discarded amount: full amount administered	

Important Billing Instructions¹:

MiroDerm is listed on the Medicare Part A and B Average Sales Price (ASP) File published quarterly by the Centers for Medicare and Medicaid Services (CMS)

- ASP information is published quarterly by the Centers for Medicare and Medicaid Services (CMS) in the ASP
 Medicare Part B Drug Pricing File or Not Otherwise Classified (NOC) Pricing File. Providers are encouraged to
 review the ASP Pricing files posted quarterly by CMS and listed by the HCPCS on CMS.gov
- The payment allowance limits for drugs and biologicals that are not included in the ASP Medicare Part B Drug
 Pricing File or Not Otherwise Classified (NOC) Pricing File, are based on the published Wholesale Acquisition
 Cost (WAC) or invoice pricing. In determining the payment limit based on WAC, the contractors follow the
 methodology specified in Publication. 100-4, Chapter 17, Drugs and Biologicals, for calculating the AWP, but
 substitute WAC for AWP.
- Providers should check with each Medicare Administrative Contractor and other payers to determine if an invoice is required to be submitted with the claim in Box 19.
- Providers should check with local payers regarding appropriate use of modifiers.
- JW Modifier must be reported for dates of service on or after January 1, 2023.
- JZ Modifier must be reported for dates of service on or after July 1, 2023.

Use either JW or JZ Modifiers to detail wastage or lack of wastage²:

Providers and suppliers are required to report the JW or JZ modifier on Medicare Part B drug claims for discarded drugs and biologicals. Also, providers and suppliers must document the amount of discarded drugs or biologicals in Medicare beneficiaries' medical record including:

- Date, time, and location of ulcer treated
- Name of skin substitute and how product supplied
- Approximate amount of product unit used
- Approximate amount of product unit discarded
- Reason for the wastage
- Manufacturer's serial/lot/batch or other unit identification number on graft material

The amount discarded should be billed on a separate line with the JW modifier. The unit field should reflect the amount of tissue discarded. The modifier is not required if no discarded portion is being billed to any payer.

Please refer to: https://www.cms.gov/Outreach-and-Education/Medicare-Learning-Network-MLN/MLNMattersArticles/Downloads/MM9603.pdf and the FAQs at https://www.cms.gov/Medicare/Medicare-Fee-for-Service-Payment/HospitalOutpatientPPS/Downloads/JWModifier-FAQs.pdf.

Use modifier JZ on billing claims to attest there was no discarded amount from the single-dose vial or single-use package that is normally paid under Part B. Reference (IOM 100-4 Chapter 17, Sections 40-40.1) or page 621 of the CMS CY2023 OPPS/ASC Final Rule: https://public-inspection.federalregister. gov/2022-23918.pdf.

^{1. &}lt;a href="https://www.cms.gov/medicare/payment/part-b-drugs/asp-pricing-files">https://www.cms.gov/medicare/payment/part-b-drugs/asp-pricing-files

Please refer to: https://www.cms.gov/Outreach-and-Education/Medicare-Learning-Network-MLN/MLNMattersArticles/Downloads/MM9603.pdf and the FAQs at https://www.cms.gov/Outreach-and-Education/Medicare-Learning-Network-MLN/MLNMattersArticles/Downloads/JMM0616 and Use modifier JZ on billing claims to attest there was no discarded amount from the single-use package that is normally paid under Part B. Reference (IOM 100-4 Chapter 17, Sections 40-40.1) or page 621 of the CMS CY2023 OPPS/ASC Final Rule: https://public-inspection.federalregister.gov/2022-23918.pdf. No changes were made to this instruction in Federal Register updates for CY2024 or CY2025. Please check with your specific MAC for additional instructions.

Place-of-Service Codes¹

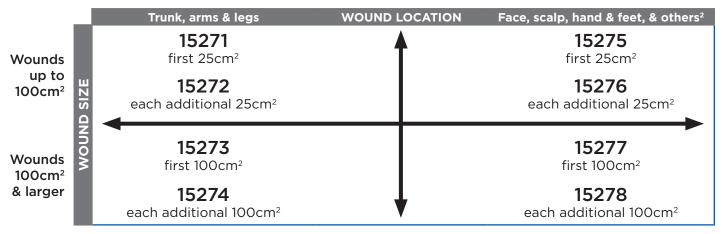
Place-of-service codes are 2-digit numbers included on health care professional claims to indicate the setting in which a service was provided. The Centers for Medicare and Medicaid Services (CMS) maintain place-of-service codes used throughout the health care industry. Listed below are place-of-service and descriptions that typically apply to our products. These codes should be used on professional claims to specify the entity where service(s) were rendered. Check with individual payors for reimbursement policies regarding these codes.

PLACE-OF-SERVICE CODE	PLACE-OF-SERVICE NAME	PLACE-OF-SERVICE DESCRIPTION
11	Office	Location other than hospital, skilled nursing facility (SNF), military treatment facility, community health center, State or Local public health clinic, or intermediate care facility (ICF), where the health professional routinely provides health examinations, diagnosis, and treatment of illness or injury on an ambulatory basis.
12	Home	Location other than a hospital or other facility, where the patient receives care in a private residence.
21	Inpatient Hospital	A facility, other than psychiatric, which primarily provides diagnostic, therapeutic (both surgical and nonsurgical) and rehabilitation services by, or under, the supervision of physicians to patients admitted for a variety of medical conditions.
22	Outpatient Hospital	A portion of a hospital's main campus which provides diagnostic, therapeutic (both surgical and nonsurgical), and rehabilitation services to sick or injured persons who do not require hospitalization or institutionalization. Description change effective January 1, 2016.
24	Ambulatory Surgical Center	A freestanding facility, other than a physician's office, where surgical and diagnostic services are provided on an ambulatory basis.
31	Skilled Nursing Facility	A facility which primarily provides inpatient skilled nursing care and related services to patients who require medical, nursing, or rehabilitative services but does not provide the level of care or treatment available in a hospital.
32	Nursing Facility	A facility which primarily provides to residents skilled nursing care and related services for the rehabilitation of injured, disabled, or sick persons, or, on a regular basis, health-related care services above the level of custodial care to other than individuals with intellectual disabilities.

^{1.} https://www.cms.gov/medicare/coding-billing/place-of-service-codes/code-sets

CPT® Coding

The Current Procedural Terminology (CPT) code set describes medical, surgical, and diagnostic services and is designed to communicate uniform information about medical services and procedures among physicians, coders, patients, accreditation organizations, and payors for administrative, financial, and analytical purposes.



CPT®	DESCRIPTIONS FOR APPLICATION OF SKIN SUBSTITUTES
15271	Application of skin substitute graft to trunk, arms, legs, total wound surface up to 100 sq. cm; first 25 sq. cm or less wound surface area.
+15272	Each additional 25 sq. cm up to 100 sq. cm wound surface area, or part thereof. List separately in addition to code 15721 for primary procedure.
15273	Application of skin substitute graft to trunk, arms, legs, total wound surface area greater than or equal to 100 sq. cm; first 100 sq. cm wound surface area, 1% of body area of infants and children.
+15274	Each additional 100 sq. cm wound surface area or part thereof, or each additional 1% of body area of infants and children or part thereof. List separately in addition to code 15273 for primary procedure.
15275	Application of skin substitute graft to face, scalp eyelids, mouth, neck, orbits, genitalia, hands feet, and/or multiple digits, total wound surface area up to 100 sq. cm; first 25cm or less wound surface area.
+15276	Each additional 25 sq. cm wound surface area, or part thereof. List separately in addition to code 15275 for primary procedure.
15277	Application of skin substitute graft to face, scalp, eyelids, mouth, neck, ears, orbits, genitalia, hands, feet, and/or multiple digits, total wound surface area greater than or equal to 100 sq. cm, first 100 sq. cm wound surface area, or 1% of body area of infants.
+15278	Each additional 100 sq. cm wound surface area, part thereof. List separately in addition to code 15277 for primary procedure.

CPT® Codes 15271-15278:

- Billing Units = 1 unit per service for CPT® 15271, 15273, 15275 and 15277 (daily limitations apply)
- Add-on codes 15272, 15274, 15276 and 15278 are billed as 1 unit for each additional amount of graft material as specified; either each additional 25cm² or 100cm² applied.

Add-on Codes: The + symbol signifies an add-on code. An add-on code cannot be used alone but must be billed with the initial code above it. Please check the CPT® 2024 coding book for further instructions.

2025 Physician Services - Medicare Payment

CHRONI	CHRONIC WOUNDS REPAIR		HOSPITAL/ASC
CPT Code ¹	Description	Medicare National Avg²	Medicare National Avg ²
15271	Application of skin substitute graft to trunk, arms, legs, total wound surface up to 100 sq. cm; first 25 sq. cm or less wound surface area	\$148.49	\$81.52
+15272	each additional 25 sq. cm wound surface area, or part thereof (List separately in addition to code for primary procedure)	\$23.62	\$16.18
15273	Application of skin substitute graft to trunk, arms, legs, total wound surface area greater than or equal to 100 sq. cm; first 100 sq. cm wound surface area, or 1% of body area of infants and children	\$295.03	\$187.31
+15274	each additional 100 sq. cm wound surface area, or part thereof, or each additional 1% of body area of infants and children, or part thereof (List separately in addition to code for primary procedure)	\$76.99	\$42.38
15275	Application of skin substitute graft to face, scalp, eyelids, mouth, neck, ears, orbits, genitalia, hands, feet, and/or multiple digits, total wound surface area up to 100 sq. cm; first 25 sq. cm or less wound surface area	\$153.99	\$90.58
+15276	each additional 25 sq. cm wound surface area, or part thereof (List separately in additional to code for primary procedure)	\$31.70	\$23.94
15277	Application of skin substitute graft to face, scalp, eyelids, mouth, neck, ears, orbits, genitalia, hands, feet, and/or multiple digits, total wound surface area greater than or equal to 100 sq. cm; first 100 sq. cm wound surface area, or 1% of body area of infants and children	\$329.97	\$215.77
+15278	each additional 100 sq. cm wound surface area, or part thereof, or each additional 1% of body area of infants and children, or part there of (List separately in addition to code for primary procedure)	\$91.23	\$54.02

Coinsurance/Deductibles: As with all products and services paid for under Medicare Parts A & B. Medicare will reimburse 80 percent of the allowable amount. The patient, or secondary/supplemental plan, is responsible for the remaining, 20 percent coinsurance amount. The appropriate annual deductions also apply.

Sequestration³: Sequestration is a mechanism for 3 budget enforcement rules created by the 1) Statutory Pay-As-You-Go Act of 2010 (Statutory PAYGO; P.L. 111-139), 2) the Budget Control Act of 2011 (BCA; P.L. 112-25), and 3) the Fiscal Responsibility Act of 2023 (FRA; P.L. 118-5). In 2024, only the BCA sequester has been triggered and is in effect at 2%. Under the BCA, the sequestration of mandatory spending was originally scheduled to occur in FY2013 through FY2021. However, subsequent legislation extended sequestration for mandatory spending through FY2031 and the sequestration of only Medicare benefit payments spending through FY2032. (The sequestration to Medicare was temporarily suspended from May 1, 2020, through March 30, 2022, and was limited to 1% from April 1, 2022, through June 30, 2022.) The Statutory PAYGO sequester applies to mandatory funding, is current law, and can be triggered if associated budget enforcement rules are broken. Due to the potential impact of the American Rescue Plan Act of 2021 (ARPA; P.L. 117-2) on deficits, sequestration under PAYGO was expected to be triggered in early 2022. However, the Protecting Medicare and American Farmers from Sequester Cuts Act (P.L. 117-71) deferred the impact of ARPA to 2023. Subsequently, the Consolidated Appropriations Act, 2023, deferred the impact of ARPA and other legislation estimated to impact the deficit under PAYGO. Without related congressional action, reductions to Medicare under PAYGO could occur in 2025. The FRA sequester applies to discretionary funding, is current law, and can be triggered if associated budget enforcement rules are broken (and Congress does not take action to change or waive this rule).

Geographic Practice Cost Index (GPCI)⁴: The Medicare physician fee schedule amounts are adjusted to reflect the variation in practice costs from area to area. A GPCI has been established for every Medicare payment locality based on the RVUs for work, practice expense, and malpractice. The GPCIs are applied in the calculation of a fee schedule payment amount by multiplying the RVU for each component times the GPCI for that component.

Non-Physician Practioners (NPP)⁵: CMS reimburses NPP professional services at 80% of the lesser of the actual charge or 85% of the amount a physician gets under the https://www.cms.gov/medicare/physician-fee-schedule/search/overview" Physician Fee Schedule(PFS) when furnished outside a hospital or SNF setting. CMS reimburses https://www.cms.gov/medicare/payment/fee-schedules/physician-fee-schedule/advanced-practice-providers/incident-services-supplies "incident to" services provided by auxiliary personnel (outside a hospital or SNF setting) at 85% of the amount a physician gets under the PFS.

^{1. 2025} AMA CPT* Professional. Procedure coding should be based upon medical necessity, procedures and supplies provided to the patient. Coding and reimbursement information is provided for educational purposes and does not assure coverage of the specific item or service in a given case. Reprise Biomedical makes no guarantee of coverage or reimbursement of fees. These payment rates are nationally unadjusted average amounts and do not account for differences in payment due to geographic variation. Contact your Medicare Administrative Contractor (MAC) or CMS for specific information as payment rates listed are subject to change. To the extent that you submit cost information to Medicare, Medicaid or any other reimbursement program to support claims for services or items, you are obligate to accurately report the actual price paid for such items, including any subsequent adjustments. CPT five-digit numeric codes, descriptions, and numeric modifiers only are Copyright AMA.

^{2.} CY2025 MPFS: https://www.cms.gov/medicare/payment/fee-schedules/physician/federal-regulation-notices/cms-1807-f CMS CY2025 MPFS Conversion factor = \$32,3465

^{3.} https://sgp.fas.org/crs/misc/R45106.pdf

^{4. &}lt;a href="https://www.cms.gov/medicare/medicare-fee-for-service-payment/physicianfeesched/downloads/geographic_adjustment_of_medicare_physician_payments_july2012.pdf and https://www.cms.gov/medicare/physician-fee-schedule/search/documentation</p>

^{5.} Add a footnote to this with this link: https://www.cms.gov/Regulations-and-Guidance/payment/fee-schedules/physician-fee-schedule/advanced-practice-nonphysician-practitioners/advanced-practice-registered-nurses-aprn; https://www.cms.gov/Regulations-and-Guidance/Guidance/Manuals/Downloads/bp102c15.pdf#page%3D149 Section 200 of the Medicare Benefit Policy Manual, Chapter 15

2025 Hospital Outpatient/ASC - Medicare

CMS Skin Substitutes Coding and Payment¹-High-Cost

In 2024, the payment for skin substitute products that do not qualify for hospital OPPS pass-through status are packaged into the OPPS payment for the associated skin substitute application procedure. This policy is also implemented in the ASC payment system. Skin substitute products are divided into two groups: 1) high-cost skin substitute products and 2) low-cost skin substitute products for packaging purposes. ASCs should not separately bill for packaged skin substitutes (SC PI=NI) since packed codes are not reportable under the ASC payment system.

A2031, MiroDry, per square centimeter	Effective January 1st 2023, CMS assigns HCPCS "A2XXX" codes to the high-cost skin substitute category. CMS made no CY2024/CY2025 updates to this direction. Therefore, MiroDry is reimbursed in the high-cost category.
Q4175 MiroDerm per square centimeter	Q4175 MiroDerm per square centimeter substitute is assigned to the high-cost category ³

CHRON	CHRONIC WOUND REPAIR		OUTPATIENT HOSPITAL		ASC	
CPT Code ¹	Description	APC	OPSI Code	Medicare National Avg Allowance⁴	Payment Indicator	Medicare National Avg Allowance⁵
15271	Application of skin substitute graft to trunk, arms, legs, total wound surface up to 100 sq. cm; first 25 sq. cm or less wound surface area	5054	Т	\$1829.23	G2	\$981.09
+15272	each additional 25 sq. cm wound surface area, or part thereof (List separately in addition to code for primary procedure)	-	N	\$0.00	N1	\$0.00
15273	Application of skin substitute graft to trunk, arms, legs, total wound surface area greater than or equal to 100 sq. cm; first 100 sq. cm wound surface area, or 1% of body area of infants and children	5055	Т	\$3660.97	G2	\$1957.33
+15274	each additional 100 sq. cm wound surface area, or part thereof, or each additional 1% of body area of infants and children, or part thereof (List separately in addition to code for primary procedure)	-	N	\$0.00	N1	\$0.00
15275	Application of skin substitute graft to face, scalp, eyelids, mouth, neck, ears, orbits, genitalia, hands, feet, and/or multiple digits, total wound surface area up to 100 sq. cm; first 25 sq. cm or less wound surface area	5054	Т	\$1829.23	G2	\$981.09
+15276	each additional 25 sq. cm wound surface area, or part thereof (List separately in additional to code for primary procedure)	-	N	\$0.00	N1	\$0.00
15277	Application of skin substitute graft to face, scalp, eyelids, mouth, neck, ears, orbits, genitalia, hands, feet, and/or multiple digits, total wound surface area greater than or equal to 100 sq. cm; first 100 sq. cm wound surface area, or 1% of body area of infants and children	5054	Т	\$1829.23	G2	\$981.09
+15278	each additional 100 sq. cm wound surface area, or part thereof, or each additional 1% of body area of infants and children, or part there of (List separately in addition to code for primary procedure)	-	N	\$0.00	N1	\$0.00

OPSI (Outpatient Payment Status Indicator)

Code T - Significant Procedure, Multiple Reduction Applies

Payment Indicator G2 - Non-office-based surgical procedure; payment based on OPPS relative payment weight

Payment Indicator N1 - Packaged service/item, no separate payment made

APC #5053 - Level III Skin Procedures; APC #5054 - Level IV Skin Procedures; APC #5055 - Level V Skin Procedures

Coinsurance/Deductibles: As with all products and services paid for under Medicare Parts A & B. Medicare will reimburse 80 percent of the allowable amount. The patient, or secondary/supplemental plan, is responsible for the remaining, 20 percent coinsurance amount. The appropriate annual deductions also apply.

Sequestration²: Sequestration is a mechanism for 3 budget enforcement rules created by the 1) Statutory Pay-As-You-Go Act of 2010 (Statutory PAYGO; P.L. 111-139), 2) the Budget Control Act of 2011 (BCA; P.L. 112-25), and 3) the Fiscal Responsibility Act of 2023 (FRA; P.L. 118-5). In 2024, only the BCA sequester has been triggered and is in effect at 2%. Under the BCA, the sequestration of mandatory spending was originally scheduled to occur in FY2013 through FY2021. However, subsequent legislation extended sequestration for mandatory spending through FY2031 and the sequestration of only Medicare benefit payments spending through FY2032. (The sequestration to Medicare was temporarily suspended from May 1, 2020, through March 30, 2022, and was limited to 1% from April 1, 2022, through June 30, 2022.) The Statutory PAYGO sequester applies to mandatory funding, is current law, and can be triggered if associated budget enforcement rules are broken. Due to the potential impact of the American Rescue Plan Act of 2021 (ARPA; P.L. 117-2) on deficits, sequestration under PAYGO was expected to be triggered in early 2022. However, the Protecting Medicare and American Farmers from Sequester Cuts Act (P.L. 117-71) deferred the impact of ARPA to 2023. Subsequently, the Consolidated Appropriations Act, 2023, deferred the impact of ARPA and other legislation estimated to impact the deficit under PAYGO. Without related congressional action, reductions to Medicare under PAYGO could occur in 2025. The FRA sequester applies to discretionary funding, is current law, and can be triggered if associated budget enforcement rules are broken (and Congress does not take action to change or waive this rule).

Geographic Practice Cost Index (GPCI)³: The Medicare physician fee schedule amounts are adjusted to reflect the variation in practice costs from area to area. A GPCI has been established for every Medicare payment locality based on the RVUs for work, practice expense, and malpractice. The GPCIs are applied in the calculation of a fee schedule payment amount by multiplying the RVU for each component times the GPCI for that component.

2. https://sgp.fas.org/crs/misc/R45106.pdf

4. CMS Addendum A OPPS APCs for 2025: https://www.cms.gov/license/ama?file=/files/zip/2025-nfrm-opps-addenda.zip

^{1. 2025} AMA CPT* Professional. Procedure coding should be based upon medical necessity, procedures and supplies provided to the patient. Coding and reimbursement information is provided for educational purposes and does not assure coverage of the specific item or service in a given case. Reprise Biomedical makes no guarantee of coverage or reimbursement of fees. These payment rates are nationally unadjusted average amounts and do not account for differences in payment due to geographic variation. Contact your Medicare Administrative Contractor (MAC) or CMS for specific information as payment rates listed are subject to change. To the extent that you submit cost information to Medicare, Medicaid or any other reimbursement program to support claims for services or items, you are obligate to accurately report the actual price paid for such items, including any subsequent adjustments. CPT five-digit numeric codes, descriptions, and numeric modifiers only are Copyright AMA.

^{3. 2025} Hospital Outpatient/ASC Rule and OPPS/ASC Addenda: https://www.cms.gov/medicare/payment/prospective-payment-systems/ambulatory-surgical-center-asc/asc-regulations-and-notices/cms-1809-fc

^{5.} CMS Addendum AA = ASC Covered Surgical Procedures for CY 2025 https://www.cms.gov/license/ama?file=/files/zip/january-2025-asc-approved-hcpcs-code-and-payment-rates.zip

Hospital Inpatient — Medicare 2025 Medicare MS-DRG* National Averages

MS-DRG	MS-DRG DESCRIPTION	MEDICARE NATIONAL AVERAGE PAYMENT ¹
570	SKIN DEBRIDEMENT W MCC	\$20,932
571	SKIN DEBRIDEMENT W CC	\$11,645
572	SKIN DEBRIDEMENT W/O CC/MCC	\$7,943
573	SKIN GRAFT FOR SKIN ULCER OR CELLULITIS W MCC	\$42,817
574	SKIN GRAFT FOR SKIN ULCER OR CELLULITIS W CC	\$24,091
575	SKIN GRAFT FOR SKIN ULCER OR CELLULITIS W/O CC/MCC	\$13,882
576	SKIN GRAFT EXC FOR SKIN ULCER OR CELLULITIS W MCC	\$37,512
577	SKIN GRAFT EXC FOR SKIN ULCER OR CELLULITIS W CC	\$18,514
578	SKIN GRAFT EXC FOR SKIN ULCER OR CELLULITIS W/O CC/MCC	\$11,753
579	OTHER SKIN, SUBCUT TISS & BREAST PROC W MCC	\$22,662
580	OTHER SKIN, SUBCUT TISS & BREAST PROC W CC	\$12,328
581	OTHER SKIN, SUBCUT TISS & BREAST PROC W/O CC/MCC	\$10,007
622	SKIN GRAFTS & WOUND DEBRID FOR ENDOC, NUTRIT & METAB DIS W MCC	\$26,033
623	SKIN GRAFTS & WOUND DEBRID FOR ENDOC, NUTRIT & METAB DIS W CC	\$13,309
624	SKIN GRAFTS & WOUND DEBRID FOR ENDOC, NUTRIT & METAB DIS W/O CC/MCC	\$6,975
628	OTHER ENDOCRINE, NUTRIT & METAB O.R. PROC W MCC	\$27,440
629	OTHER ENDOCRINE, NUTRIT & METAB O.R. PROC W CC	\$15,646
630	OTHER ENDOCRINE, NUTRIT & METAB O.R. PROC W/O CC/MCC	\$9,743
904	SKIN GRAFTS FOR INJURIES W CC/MCC	\$26,846
905	SKIN GRAFTS FOR INJURIES W/O CC/MCC	\$11,460
907	OTHER O.R. PROCEDURES FOR INJURIES W MCC	\$27,699
908	OTHER O.R. PROCEDURES FOR INJURIES W CC	\$14,025
909	OTHER O.R. PROCEDURES FOR INJURIES W/O CC/MCC	\$8,818

The table above provides potential MS-DRGs assignments for hospitals when applying MiroDerm. These are 2025 Medicare average rates and are provided as a benchmark. Rates nationwide can vary widely.

*Medicare reimburses hospital inpatient stays based on the Medicare Severity Diagnosis Related Group (MS-DRG) system. MS-DRGs represent a consolidated prospective payment for all services provided by the hospital during hospitalization, based on submitted claims data. With limited exceptions, the MS-DRG payment is inclusive of all services, products, and resources, regardless of the final cost to the hospital. Medicare and many private payors use the MS-DRG based system to reimburse facilities for inpatient services.

ICD-10-Procedure Code: OHR5XK3

Replacement of Chest Skin with Nonautologous Tissue Substitute, Full Thickness, External Approach

Effective 10.01.2022, replacement of skin using a porcine liver-derived skin substitute procedure code moved from the new technology section to the medical and surgical section, skin and breast body system with device value as nonautologous.

Hospital inpatient procedures are billed with ICD-10 CM Procedure Codes. These procedure codes are mapped to specific Diagnosis Related Groups (DRGs) for payment. Private payor claims processing protocol can vary, but often use the same DRG Grouper as Medicare resulting in assignment to the same DRGs.

NOTE: This list is not all inclusive. These procedure codes, when combined with appropriate ICD-10 CM diagnosis codes get mapped to DRGs for payment. The ICD-10 PCS system provides greater granularity in the reporting of procedures.

- 1. CMS Hospital Inpatient Final Rule is effective October 1 2024 September 30 2025. Beginning in FFY 2023, CMS adopted a permanent 10% cap on reductions to a MS-DRG's relative weight in a given year compared to the weight in the prior year, implemented in a budget-neutral manner. The cap will be applied regardless of the reason for the decrease and implemented in a budget-neutral manner.
- Medicare Program; Hospital Inpatient Prospective Payment Systems for Acute Care Hospitals and the Long-Term Care Hospital Prospective Payment System and Policy Changes
 and Fiscal Year 2025 Rates; Quality Programs and Medicare Promoting Interoperability Program Requirements for Eligible Hospitals and Critical Access Hospitals; Rural
 Emergency Hospital and Physician-Owned Hospital Requirements; and Provider and Supplier Disclosure of Ownership; and Medicare Disproportionate Share Hospital (DSH)
 Payments: Counting Certain Days Associated with Section 1115 Demonstrations in the Medicaid Fraction: https://public-inspection.federalregister.gov/2023-16252.pdf

ICD-10-CM¹ Diagnosis Code Examples Appendix

ICD-10 diagnosis codes identify a patient's condition, while ICD-10 procedure codes identify the services or treatments a patient receives.

^{1.} The International Classification of Diseases Clinical Modification Edition 9 is developed by the National Center for Health Statistics (NCHS). ICD-9 CM and ICD-10 CM are copyrighted by the World Health Organization. WHO is the copyright holder of ICD-10, and can grant licenses for the use of ICD-10 worldwide for commercial and non-commercial uses.

ICD-10-CM Diagnosis Examples

The ICD-10-CM codes listed below represent some of the etiology diagnosis codes commonly associated with causes of lower extremity chronic ulcers. This is not meant to be an exhaustive list. The below list of codes includes an edit to use an additional ICD-10-CM manifestation code from the L97 non-pressure chronic ulcer code series as a secondary diagnosis.

ICD-10-CM DIAGNOSES CODES	DESCRIPTION
E10.621	Type 1 diabetes mellitus with foot ulcer
E10.622	Type 1 diabetes mellitus with other skin ulcer
E11.621	Type 2 diabetes mellitus with foot ulcer
E11.622	Type 2 diabetes mellitus with other skin ulcer
E13.621	Other specified diabetes mellitus with foot ulcer
E13.621	Other specified diabetes mellitus with other skin ulcer
183.012	Varicose veins of right lower extremity with ulcer of calf
183.013	Varicose veins of right lower extremity with ulcer of ankle
183.014	Varicose veins of right lower extremity with ulcer of heel & midfoot
183.015	Varicose veins of right lower extremity with ulcer of other part of foot
183.018	Varicose veins of right lower extremity with ulcer of other part of lower leg
183.022	Varicose veins of left lower extremity with ulcer of calf
183.023	Varicose veins of left lower extremity with ulcer of ankle
183.024	Varicose veins of left lower extremity with ulcer of heel & midfoot
183.025	Varicose veins of left lower extremity with ulcer of other part of foot
183.028	Varicose veins of left lower extremity with ulcer of other part of lower leg
183.212	Varicose veins of right lower extremity with both ulcer of calf and inflammation
183.213	Varicose veins of right lower extremity with both ulcer of ankle and inflammation
183.214	Varicose veins of right lower extremity with both ulcer of heel & midfoot and inflammation
183.215	Varicose veins of right lower extremity with both ulcer of other part of foot and inflammation
183.218	Varicose veins of right lower extremity with both ulcer of other part of lower extremity and inflammation
183.222	Varicose veins of left lower extremity with both ulcer of calf and inflammation
183.223	Varicose veins of left lower extremity with both ulcer of ankle and inflammation
183.224	Varicose veins of left lower extremity with both ulcer of heel & midfoot and inflammation
183.225	Varicose veins of left lower extremity with both ulcer of other part of foot and inflammation
183.228	Varicose veins of left lower extremity with both ulcer of other part of lower extremity and inflammation
187.2	Venous Insufficiency (chronic peripheral)
187.311	Chronic venous hypertension (idiopathic) with ulcer of right lower extremity
187.312	Chronic venous hypertension (idiopathic) with ulcer of left lower extremity
187.313	Chronic venous hypertension (idiopathic) with ulcer of bilateral lower extremity
187.331	Chronic venous hypertension (idiopathic) with ulcer and inflammation of right lower extremity
187.332	Chronic venous hypertension (idiopathic) with ulcer and inflammation of left lower extremity
187.333	Chronic venous hypertension (idiopathic) with ulcer and inflammation of bilateral lower extremity

ICD-10-CM Diagnosis Examples

The ICD-10-CM codes listed below are specific manifestation diagnosis codes commonly associated with non-pressure chronic ulcers of the lower extremity. This is not meant to be an exhaustive list.

ICD-10-CM DIAGNOSES CODES	DESCRIPTION
L97	Series Non-Pressure Chronic Ulcer of Lower limb
L97.211	Non-Pressure Chronic Ulcer of Right calf limited to breakdown of skin
L97.212	Non-Pressure Chronic Ulcer of Right calf with fat layer exposed
L97.213	Non-Pressure Chronic Ulcer of Right calf with necrosis of muscle
L97.214	Non-Pressure Chronic Ulcer of Right calf with necrosis of bone
L97.221	Non-Pressure Chronic Ulcer of Left calf limited to breakdown of skin
L97.222	Non-Pressure Chronic Ulcer of Left calf with fat layer exposed
L97.223	Non-Pressure Chronic Ulcer of Left calf with necrosis of muscle
L97.224	Non-Pressure Chronic Ulcer of Left calf with necrosis of bone
L97.311	Non-Pressure Chronic Ulcer of Right ankle limited to breakdown of skin
L97.312	Non-Pressure Chronic Ulcer of Right ankle with fat layer exposed
L97.313	Non-Pressure Chronic Ulcer of Right ankle with necrosis of muscle
L97.314	Non-Pressure Chronic Ulcer of Right ankle with necrosis of bone
L97.321	Non-Pressure Chronic Ulcer of Left ankle limited to breakdown of skin
L97.322	Non-Pressure Chronic Ulcer of Left ankle with fat layer exposed
L97.323	Non-Pressure Chronic Ulcer of Left ankle with necrosis of muscle
L97.324	Non-Pressure Chronic Ulcer of Left ankle with necrosis of bone
L97.411	Non-Pressure Chronic Ulcer of Right heel & midfoot limited to breakdown of skin
L97.412	Non-Pressure Chronic Ulcer of Right heel & midfoot with fat layer exposed
L97.413	Non-Pressure Chronic Ulcer of Right heel & midfoot with necrosis of muscle
L97.414	Non-Pressure Chronic Ulcer of Right heel & midfoot with necrosis of bone
L97.421	Non-Pressure Chronic Ulcer of Left heel & midfoot limited to breakdown of skin
L97.422	Non-Pressure Chronic Ulcer of Left heel & midfoot with fat layer exposed
L97.423	Non-Pressure Chronic Ulcer of Left heel & midfoot with necrosis of muscle
L97.424	Non-Pressure Chronic Ulcer of Left heel & midfoot with necrosis of bone
L97.511	Non-Pressure Chronic Ulcer of Other part of right foot limited to breakdown of skin
L97.512	Non-Pressure Chronic Ulcer of Other part of right foot with fat layer exposed
L97.513	Non-Pressure Chronic Ulcer of Other part of right foot with necrosis of muscle
L97.514	Non-Pressure Chronic Ulcer of Other part of right foot with necrosis of bone
L97.521	Non-Pressure Chronic Ulcer of Other part of left foot limited to breakdown of skin
L97.522	Non-Pressure Chronic Ulcer of Other part of left foot with fat layer exposed
L97.523	Non-Pressure Chronic Ulcer of Other part of left foot with necrosis of muscle
L97.524	Non-Pressure Chronic Ulcer of Other part of left foot with necrosis of bone
L97.811	Non-Pressure Chronic Ulcer of Other part of right lower leg limited to breakdown of skin
L97.812	Non-Pressure Chronic Ulcer of Other part of right lower leg with fat layer exposed
L97.813	Non-Pressure Chronic Ulcer of Other part of right lower leg with necrosis of muscle
L97.814	Non-Pressure Chronic Ulcer of Other part of right lower leg with necrosis of bone
L97.821	Non-Pressure Chronic Ulcer of Other part of left lower leg limited to breakdown of skin
L97.822	Non-Pressure Chronic Ulcer of Other part of left lower leg with fat layer exposed
L97.823	Non-Pressure Chronic Ulcer of Other part of left lower leg with necrosis of muscle
L97.824	Non-Pressure Chronic Ulcer of Other part of left lower leg with necrosis of bone

Sample Letter of Medical Necessity*

[INSTRUCTION: PICK APPROPRIATE PRODUCT]

Date Insurer Name Insurer Address City, State, Zip Code

RE: Medical Necessity for MiroDry Wound Matrix or MiroDerm Biologic Wound Matrix Patient's Name:
Policy Number:
Group Number:

Date of Birth:

Dear [Insurance Contact Name]:

I am writing to notify you of my intent to treat Mr./Ms. [Patient's Name] with [Insert product name]. It is used for the management of wounds, including:

[FOR MIRODRY WOUND MATRIX:] partial and full-thickness wounds; pressure ulcers; venous ulcers; chronic vascular ulcers; diabetic ulcers; tunneled, undermined wounds; trauma wounds (abrasion, lacerations, partial thickness burns, skin tears); drainage wounds; and surgical wounds (donor sites/grafts, post-Mohs surgery, post-laser surgery, podiatric, wound dehiscence).

[FOR MIRODERM BIOLOGIC WOUND MATRIX:] partial and full-thickness wounds; pressure ulcers; venous ulcers; chronic vascular ulcers; diabetic ulcers; tunneled, undermined wounds; trauma wounds (abrasion, lacerations, second-degree burns, skin tears); drainage wounds; and surgical wounds (donor sites/grafts, post-Mohs surgery, post-laser surgery, podiatric, wound dehiscence).

The patient's medical history is as follows: [include relevant medical history]

[FOR MIRODRY WOUND MATRIX:] MiroDry is a dry, open, and porous collagen sheet matrix that provides a protective environment for wound management. MiroDry was cleared by the FDA under 510(k) K240277.

[FOR MIRODERM BIOLOGIC WOUND MATRIX:] MiroDerm is an acellular wound matrix that is derived from the highly vascularized porcine liver and was cleared by the FDA under 510(k) K143426.

[Include the following two paragraphs if the wound is an ulcer; otherwise, do not include them. Examples of wounds for which the paragraphs should NOT be used include pilonidal wounds or wound dehiscence.] [Insert product name], being derived from porcine sources, is fully covered by National Coverage Decision (NCD) 270.5 - Porcine Skin and Gradient Dressings. The NCD's coverage criteria, which include "burns, donor sites of a homograft, and decubiti and other ulcers" (emphasis added), confirm that [Insert product name] qualifies as a covered product for ulcers, as experienced by [insert patient's name], when deemed reasonable and necessary by the treating provider. I believe the use of [Insert product name] is reasonable, necessary, and beneficial for [insert patient's name].

Reprise Biomedical manufactures **[insert product name]** and is a medical device company focused on the development of clinically relevant biologic medical devices for the surgical and wound care space.

My patient has not responded to conservative care for [time frame] and has not responded to more advanced therapy including [product name(s) & type(s) of products]. More aggressive treatment is medically necessary to prevent further damage and [list risk(s) of non-closure]. I believe my patient will benefit from treatment with [Insert product name].

I have enclosed information regarding the clinical utility of [Insert product name].

Please feel free to contact me if additional information is required to process my request for coverage.

Sincerely,

[Name] [Contact info]

^{*}This sample letter contains content for both MiroDry and MiroDerm. Please use the appropriate provided language for the desired product.

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