

**PATIENT INFORMATION**

Order date: \_\_\_\_\_

Patient Name: \_\_\_\_\_  
Last First MI DOB: \_\_\_\_\_

Home Address: \_\_\_\_\_ Phone: \_\_\_\_\_

City: \_\_\_\_\_ State: \_\_\_\_\_ Zip: \_\_\_\_\_ Cell Phone: \_\_\_\_\_

**REFERRAL INFORMATION**

Referral Facility: \_\_\_\_\_ Contact Name: \_\_\_\_\_

Phone #: \_\_\_\_\_

**PLEASE SEND PATIENT DEMOGRAPHICS/INSURANCE INFORMATION ALONG WITH CHART NOTES**

**DELIVERY INFORMATION**

Requested Delivery Date: \_\_\_\_\_ Requested Delivery Time: \_\_\_\_\_

Delivery Location:  Home - SAME ADDRESS AS ABOVE  Other: \_\_\_\_\_

Hospital/Facility: \_\_\_\_\_

Room Number: \_\_\_\_\_

**PRESCRIPTION, ATTESTATION AND TREATING PRESCRIBER'S INFORMATION**

This form is required unless a separate detailed written order for NPWT is provided. Prescriber must clearly document in the patient's medical record that other modalities have been tried, or clearly document why other modalities are being ruled out.

Diagnosis Code ICD-10. Write in complete code(s): \_\_\_\_\_

**PHYSICIAN'S ORDER**

I prescribe a Negative Pressure Wound Therapy Pump (E2402), and up to 15 dressing kits (A6550) per wound per month and up to 10 canisters (A7000) per month.

Pressure Setting: \_\_\_\_\_  Continuously  Intermittently Frequency of Dressing Changes: \_\_\_\_\_

For the following wound type:  Surgical  Dehisced  Traumatic  Pressure Ulcer  Venous/Arterial Ulcer  
 Neuropathic/Diabetic Ulcer  Chronic Mixed Etiology (≥ 30 Days)

Wound Location: \_\_\_\_\_

Goal of NPWT:  Assist granulation tissue formation  Delayed Primary Closure  Flap/Graft

Length of Need (Anticipated):  1 Month  2 Months  3 Months  4 Months (Medicare allows 4 months with wound improvement)

Other: \_\_\_\_\_

Dressing Kits			
Foam:	<input type="checkbox"/> Small	<input type="checkbox"/> Medium	<input type="checkbox"/> Large
Gauze:		<input type="checkbox"/> Medium	<input type="checkbox"/> Large

Miscellaneous Supplies

**PRESCRIBER INFORMATION**

Prescriber Name: \_\_\_\_\_

Address: \_\_\_\_\_ City: \_\_\_\_\_ ST: \_\_\_\_\_ Zip: \_\_\_\_\_

Phone: \_\_\_\_\_ Fax: \_\_\_\_\_

**Prescriber Only to Complete—Original Signature and Date Required. No Stamps**

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

NPI #: \_\_\_\_\_

By signing and dating, I attest that I am prescribing Negative Pressure Wound Therapy (NPWT) as medically necessary and all other applicable treatments have been tried or considered and ruled out. I have read and understand all safety information and other instructions for use included with the NPWT product. I also understand NPWT contraindications. Additionally, I have reviewed the information provided in this form and attest to its accuracy.

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_\_

**PATIENT FOLLOW-UP CARE**

Home Health Agency Name: \_\_\_\_\_ Phone: \_\_\_\_\_

Wound Clinic Name: \_\_\_\_\_ Phone: \_\_\_\_\_

**COMMON ICD-10 CODES FOR NEGATIVE PRESSURE WOUND THERAPY**

Since NPWT is not diagnosis-driven, there is not a defined set of codes that must be used with this equipment. There are many other ICD-10 codes for which Negative Pressure Wound Therapy can be used. This is simply a short list of commonly used codes. Presence of an ICD-10 code alone does not guarantee coverage of a NPWT device.

**A specific ICD-10 code must be provided either on page 1 or in the patient's chart notes. Please list the appropriate qualifying diagnosis and write in the code. Ranges will not be accepted.**

ICD-10	Description	ICD-10	Description
I83.001 – I83.229	Varicose Veins with Ulcer	L89.130 – L89.229	Pressure Ulcer of Lower Back, Hip
I87.2	Venous Insufficiency (Chronic) (Peripheral)	L89.300 – L89.319	Pressure Ulcer of Buttock
L03.90	Cellulitis, Unspecified	L89.500 – L89.629	Pressure Ulcer of Lower Limb
L03.115 – L04.3	Cellulitis of Limb	T81.30XA – T81.30XS	Disruption of Wound, Unspecified
L05.01	Pilonidal Cyst with Abscess	T81.31XA – T81.31XS	Disruption of External Operation (Surgical) Wound
L89.004 – L89.894	Pressure Ulcer, Stage IV		

**WOUND TYPE (Complete in full OR fax wound history documentation)**

- Pressure Ulcer:**     Stage III     Stage IV
- Is patient being turned/positioned?  YES     NO
- Has a group 2 or 3 surface been used for ulcer located on the posterior trunk or pelvis?  YES     NO
- Are moisture and/or incontinence being managed?  YES     NO
- Is pressure ulcer greater than 30 days?  YES     NO
- Diabetic Ulcer/Neuropathic Ulcer:**
- Has a reduction of pressure on the foot ulcer been accomplished with appropriate modalities?  YES     NO
- Venous Stasis Ulcer/Venous Insufficiency:** Are compression bandages and/or garments being consistently applied?  YES     NO
- Is elevation/ambulation being encouraged?  YES     NO
- Arterial Ulcer/Arterial Insufficiency:** Is pressure over the wound being relieved?  YES     NO
- Surgical:** Wound surgically created and not represented by descriptions above?  YES     NO

Description of surgical procedure: \_\_\_\_\_ Date of surgical procedure involving wound: \_\_\_\_\_

**Chronic Ulcer of Mixed Etiology** (describe): \_\_\_\_\_

**Other Wound Type** (describe): \_\_\_\_\_

**CLINICAL WOUND INFORMATION (please submit supporting documentation)**

**WOUND #1**

- Was NPWT utilized within the last 90 days?  YES     NO
- If YES, date initiated: \_\_\_\_\_
- Is the patient's nutritional status compromised?  YES     NO
- If YES, please attach nutritional plan.
- Is osteomyelitis present in the wound?  YES     NO
- If YES, treated with: \_\_\_\_\_
- Is malignancy present in the wound?  YES     NO
- Is there a open fistula to an organ or body cavity within the vicinity of the wound?  YES     NO
- Which therapies were utilized to maintain a moist wound environment?  
 Saline/Gauze     Hydrogel     Alginate     Hydrocolloid  
 Absorptive     Other: \_\_\_\_\_

Wound location: \_\_\_\_\_ Wound Age: \_\_\_\_\_

Is wound full thickness:  YES     NO

Length: \_\_\_\_\_ cm    Width: \_\_\_\_\_ cm    Depth: \_\_\_\_\_ cm

Measurement Date: \_\_\_\_\_ Exudate Amount (daily): \_\_\_\_\_

Exudate Type: \_\_\_\_\_ Odor:  YES     NO

Please check what is exposed:

Muscle     Tendon     Bone     None

Is there tunneling?  YES     NO

If YES, Location #1 \_\_\_\_\_ cm, @ \_\_\_\_\_ o'clock

Location #2 \_\_\_\_\_ cm, @ \_\_\_\_\_ o'clock

Is there undermining?  YES     NO

If YES, Location #1 \_\_\_\_\_ cm, @ \_\_\_\_\_ o'clock

Location #2 \_\_\_\_\_ cm, @ \_\_\_\_\_ o'clock

Has a debridement been performed in the past 10 days?  YES     NO

If YES, Debridement Date: \_\_\_\_\_ Debridement Type: \_\_\_\_\_

\*Debridement needs to be attempted for the presence of necrotic tissue

**WOUND #2**

- Was NPWT utilized within the last 90 days?  YES     NO
- If YES, date initiated: \_\_\_\_\_
- Is the patient's nutritional status compromised?  YES     NO
- If YES, please attach nutritional plan.
- Is osteomyelitis present in the wound?  YES     NO
- If YES, treated with: \_\_\_\_\_
- Is malignancy present in the wound?  YES     NO
- Is there a open fistula to an organ or body cavity within the vicinity of the wound?  YES     NO
- Which therapies were utilized to maintain a moist wound environment?  
 Saline/Gauze     Hydrogel     Alginate     Hydrocolloid  
 Absorptive     Other: \_\_\_\_\_

Wound location: \_\_\_\_\_ Wound Age: \_\_\_\_\_

Is wound full thickness:  YES     NO

Length: \_\_\_\_\_ cm    Width: \_\_\_\_\_ cm    Depth: \_\_\_\_\_ cm

Measurement Date: \_\_\_\_\_ Exudate Amount (daily): \_\_\_\_\_

Exudate Type: \_\_\_\_\_ Odor:  YES     NO

Please check what is exposed:

Muscle     Tendon     Bone     None

Is there tunneling?  YES     NO

If YES, Location #1 \_\_\_\_\_ cm, @ \_\_\_\_\_ o'clock

Location #2 \_\_\_\_\_ cm, @ \_\_\_\_\_ o'clock

Is there undermining?  YES     NO

If YES, Location #1 \_\_\_\_\_ cm, @ \_\_\_\_\_ o'clock

Location #2 \_\_\_\_\_ cm, @ \_\_\_\_\_ o'clock

Has a debridement been performed in the past 10 days?  YES     NO

If YES, Debridement Date: \_\_\_\_\_ Debridement Type: \_\_\_\_\_

\*Debridement needs to be attempted for the presence of necrotic tissue