PATIENT INFORMATION

TPN Total Parenteral Nutrition Delivery Mode

(CADD&Prizm®VIP)

Model 6100 and 6101

Ambulatory

Infusion

Pumps



Deltec

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Introduction

Your doctor has recommended that you use the CADD-Prizm® pump as part of your treatment.

The CADD-Prizm® pump can be carried with you and is designed to deliver medication into your body. Your physician will prescribe your medication specifically for you. Your prescription is programmed into your pump by your clinician according to your physician's specific orders. This pump can be reprogrammed as your medication needs change.

The pump can be programmed to deliver TPN ("total parenteral nutrition") solution or other fluids at a constant rate. Delivery may "taper up," or gradually increase, at the beginning of delivery. Delivery may also "taper down," or gradually decrease, at the end of delivery. The pump stores programmed information and historical information that your clinician needs for your specific therapy.

Your clinician will instruct you on the proper use of this pump. This guide is intended to supplement those instructions. Perform only those procedures for which you have received training.

The following is a list of warnings and cautions that you should read before operating the pump. It is important that you understand and follow these warnings and cautions.

Failure to properly follow warnings, cautions, and instructions could result in damage to the pump or death or serious injury.

Warnings

- If the pump is used to deliver life-sustaining medication, an additional pump must be available.
- Use of a syringe with the CADD® Administration Set may result in UNDER-DELIVERY of medication. Syringe function can be adversely affected by variations in plunger dimension and lubricity, which can result in greater force required to move the syringe plunger. A syringe plunger will lose lubrication as it ages and, as a result, the amount of under-delivery will increase and could, on occasion, be significant.

You must regularly compare the volume remaining in the syringe to the pump's displayed values such as Reservoir Volume or Given to determine if under-delivery is occurring and, if necessary, contact your clinician.

- Do not use rechargeable NiCad or nickel metal hydride (NiMH) batteries. Do not use carbon zinc ("heavy duty") batteries. They do not provide sufficient power for the pump to operate properly.
- Always have new batteries available for replacement. If power is lost, non-delivery of drug will occur.
- There is no pump alarm to alert you that the battery has not been properly installed or has become dislodged. An improperly installed or dislodged battery could result in loss of power and non-delivery of drug.

- If the pump is dropped or hit, the battery door may become broken or damaged. Do not use the pump if the battery door is damaged because the battery will not be properly secured; this may result in loss of power or non-delivery of drug.
- Prior to starting infusion, inspect the fluid path for kinks, a closed clamp, or other upstream obstructions, and remove all air to prevent air embolism.
- Close the tubing clamp before removing the cassette from the pump to prevent unregulated gravity infusion.
- You must use a CADD® Extension Set with Anti-Siphon
 Valve or a CADD® Administration Set with either an
 integral or add on Anti-Siphon Valve to protect against
 delivery inaccuracies and unregulated gravity infusion that
 can result from an improperly attached cassette.
- For detailed instructions and warning pertaining to Medication Cassette[™] Reservoirs or CADD® Administration Sets, please refer to the instructions for use supplied with those products.
- Frozen Medication must be thawed at room temperature only. Do not heat the Medication Cassette[™] Reservoir in a microwave oven as this may damage the medication, the Medication Cassette[™] Reservoir, or cause leakage.
- Attach the cassette (the part of the Medication Cassette[™]
 Reservoir or CADD® Administration Set that attaches to
 the pump) properly. An improperly attached or detached
 cassette could result in unregulated gravity infusion or a
 reflux of blood.

- Do not prime the fluid path with the tubing connected to your catheter. This could result in over-delivery of medication or air embolism.
- Ensure that the entire fluid path is free of all air bubbles before connecting to your catheter to prevent air embolism.
- If the pump is dropped or hit, inspect the pump for damage. Do not use a pump that is damaged or is not working properly. Contact your clinician for further instructions.

Cautions

- Do not operate the pump at temperatures below + 2°C (36°F) or above 40°C (104°F).
- Do not store the pump at temperatures below -20°C (-4°F) or above 60°C (140°F). Do not store the pump with a Medication Cassette™ Reservoir or CADD® Administration Set attached.
- Do not expose the pump to humidity levels below 10% or above 90% relative humidity.
- Do not store the pump for prolonged periods with the battery installed.
- Failure to push the Power Pack cord connector or AC
 Adapter cord connector all the way forward may result in
 an intermittent connection, and the connector may dislodge, causing a loss of power and pump alarms.
- Do not immerse the pump in cleaning fluids or water or allow solution to soak into the pump, accumulate on the keypad, or enter the battery compartment, Data In/Out jack, accessory jack, or Air Detector Port area.

- Do not clean the pump with acetone, other plastic solvents, or abrasive cleaners.
- Do not use the pump in the presence of flammable anesthetics or explosive gasses.
- Use only Deltec accessories as using other brands may adversely affect the operation of the pump.

CADD-Prizm® Pump (Diagram)



Green Light

Blinks every 3 seconds when the pump is running and delivering fluid.

Amber Light

Flashing: pump is stopped; or an alarm exists. Steady: pump is inoperable, call clinician.

Display

Shows information and messages. After a short time, the display turns itself off to save power. Press any key to turn the display back on.

Description of the Keys

Starts and stops the pump and silences alarms.

LOCK Used by the clinician.

? The "Help" Key — explains what you see on the display.

ENTER Used by the clinician.

Advances you from one screen to the next and silences some alarms.

DOSE This key is not used in the TPN Delivery mode.

(OPTIONS) Used by the clinician.

Lets you answer yes to a question on the pump's display.

Lets you answer no to a question on the pump's display.

Installing a New Battery

A 9 volt battery may be used to power the pump. Even if you are using a Power Pack or AC Adapter to power the pump, you must have a 9 volt battery installed.

If 9-volt Battery Low or 9-volt Battery Depleted appears in the display, or if Battery Low appears on the main screen, you should change the battery.

Use a new 9 volt alkaline or lithium battery such as the DURACELL® Alkaline MN 1604, the EVEREADY® ENERGIZER Alkaline #522 or the ULTRALIFE® Lithium U9VL battery.

Dispose of used batteries in an environmentally safe manner, and according to any regulations which may apply.

WARNINGS:

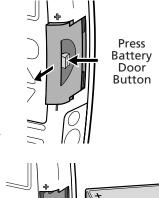
- Do not use rechargeable NiCad or nickel metal hydride (NiMH) batteries. Do not use carbon zinc ("heavy duty") batteries. They do not provide sufficient power for the pump to operate properly, which could result in death or serious injury.
- Always have new batteries available for replacement. If power is lost, non-delivery of drug will occur and, depending on the drug being administered, could result in death or serious injury.
- There is no pump alarm to alert you that the battery has not been properly installed or has become dislodged. An improperly installed or dislodged battery could result in loss of power and non-delivery of drug and, depending on the drug being administered, could result in death or serious injury.
- If the pump is dropped or hit, the battery door may become broken or damaged. DO NOT USE the pump if it

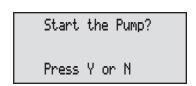
has been damaged because the battery will not be properly secured; this may result in loss of power, nondelivery of drug, and, depending on the type of drug being administered, death or serious injury.

To install a new battery:

- 1. Stop the pump by pressing (STOP) START.
- 2. When you see Stop the pump?, press .
- Press the button on the battery door and slide the battery door forward. Remove the used battery.
- 4. Match the + and markings on the new battery with the markings on the pump. Insert the battery. The pump will beep if the battery is inserted correctly.
- Replace the battery door. The pump will power up automatically.
- 6. Start the pump by pressing (STOP).
- 7. When you see Start the pump?, press .

Stop the Pump? Press Y or N





NOTE:

• If you put the battery in backwards, the display will remain

blank. Reinsert the battery, making sure to match the + and – markings.

The power up sequence will start, the pump will go through an electronic self-test, and the pump will beep at the end of the power up sequence. All of the display indicators, the software revision level, and each parameter will appear briefly.

CAUTION: Do not store the pump for prolonged periods of time with the battery installed. Battery leakage could damage the pump.

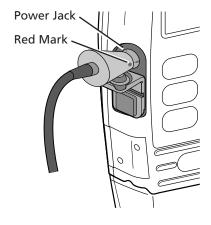
Using the Power Pack or AC Adapter

Depending on your delivery, your clinician may give you a rechargeable Power Pack or an AC Adapter to power your pump. For complete instructions, you should also read the *Instructions for Use* that come with the Power Pack or AC Adapter.

The cord from the Power Pack or AC Adapter plugs directly into the "Power" jack on the side of your pump.

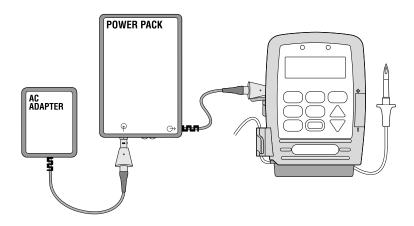
NOTE: A good 9 volt battery must also be installed in your pump as a backup. Otherwise, you won't be able to start the pump. Your Power Pack or AC Adapter may or may not have the connector shown.

- 1. Open the "Power" cover on the side of your pump.
- Line up the red mark on the Power Pack cord connector with the red mark on the pump's Power jack.
- 3. Push the connector forward until it stops. Do not twist or turn the connector. Pull lightly on the cord directly behind the flared part of the connector to make sure it is firmly attached.



CAUTION: Failure to push the connector all the way forward may result in an intermittent connection, and the connector may dislodge, causing a loss of power.

NOTE: If necessary, you can recharge the Power Pack while it is still attached to the pump. Just attach the AC Adapter connector to the Power Pack input jack. Or, if you want to operate the pump on AC power, you can attach the AC Adapter directly to the pump's Power jack.

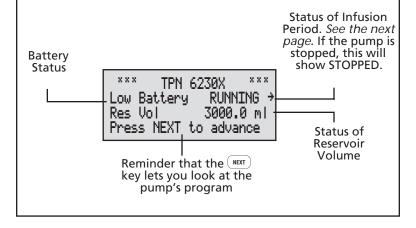


Detaching the Power Pack

- 1. Grasp the connector.
- 2. Pull the connector back using a straight, steady motion. DO NOT twist or turn the connector.

The Main Screen

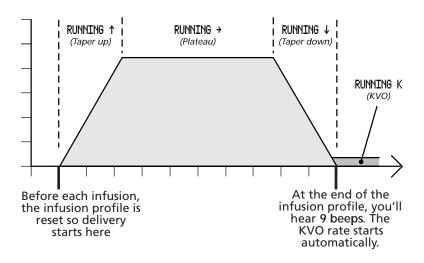
The following screen is what you will see on the pump's display most of the time. It is called the Main Screen and shows the following:



Description of the Infusion Profile

The illustration below shows what an infusion profile may look like. You can tell where you are in your infusion profile by looking at the main screen.

- During taper up, the main screen shows RUNNING †
- During the plateau rate, the main screen shows RUNNING >
- During *taper down*, the main screen shows **RUNNING** ↓
- During KVO, the main screen shows RUNNING K



Your clinician may or may not have programmed a taper up or a taper down. If there is no taper up, delivery will start at the plateau rate. If there is no taper down, you will still hear 9 beeps at the end of the infusion profile, but the rate will immediately drop from the plateau rate to the KVO rate. Then when you stop the pump, it will automatically reset to the beginning of the infusion profile.

How to Use the HELP Key (

If you have a question about a screen, press ? for more information. A description of the screen will appear.

For example, if you want information about the Reservoir Volume screen, you can press ? and this screen will appear:

This is the calculated amount of fluid left in the reservoir.

The symbol (?+) in the lower right corner means there are more help screens. Press ? again to see the next help screen.

To page through all the help screens, press ? repeatedly. This will take you through all the help screens, then back to the original screen you had questions about.

To return to the original screen you had questions about, without going through all the help screens, press (NEXT).

Starting the Pump

When you start the pump, it will review the program, and then begin delivering medication. If the pump will not start, a message should appear on the display. Refer to the Messages and Alarms section on page 26.

WARNING: Prior to starting infusion, inspect the fluid path for kinks, a closed clamp, or other upstream obstructions, and remove all air bubbles to prevent air embolism. An undetected upstream occlusion may result in under- or non-delivery of drug and, depending on the drug being administered, could result in death or serious injury. Air embolism can result in death or serious injury.

- 1. Press (STOP START).
- 2. When Start the pump? appears, press .

Starting pump... will appear. The pump will automatically review and display the preprogrammed settings.

RUNNING will appear on the main screen, the green light will blink, and fluid delivery will begin as programmed by your clinician.

Start the Pump?

Press Y or N

Starting pump...

*** TPN 6230X *** RUNNING ↑ Res Vol 3000.0 ml Press NEXT to advance

Stopping and Restarting the Pump

Stopping the pump stops delivery of medication. Whenever the pump is stopped, the amber light will blink.

- If you stop the pump *during the Infusion Profile*, delivery will stop, but will continue from the same point when you restart the pump.
- If you stop the pump during KVO delivery, the Infusion
 Profile and the Reservoir Volume will automatically be
 reset. This means that when you restart the pump, it will
 start from the beginning of the Infusion Profile.
- 1. Press (STOP START).
- 2. When Stop the Pump? appears, press .

You will see two screens if you stopped the pump during KVO delivery:

Stop the Pump?

Press Y or N

Infusion Profile has been reset

Reservoir Volume 3000.0 ml

<Resetting...>

STOPPED will appear on the main screen, and the amber light will blink.

*** TPN 6230X ***
STOPPED
Press NEXT to advance

Removing a Cassette

WARNING: Close the tubing clamp before removing the cassette from the pump to prevent unregulated infusion, which could result in death or serious injury.

To remove a cassette:

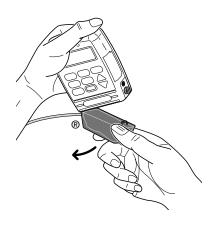
- 1. Press (STOP) to stop the pump. When Stop the pump? appears, press ...
- Stop the Pump?

 Press Y or N
- Close all tubing clamps and disconnect the tubing from your access device as instructed by your clinician.
- 3. Use a coin to unlatch the cassette. Insert the coin into the slot and turn clockwise until the latching button pops out.



NOTE: If you cannot turn the latching button, the cassette may be locked. Contact your clinician.

- 4. Remove the cassette from the pump.
- Discard the Medication
 Cassette™ Reservoir or
 CADD® Administration Set as instructed by your clinician.



Attaching a Cassette

WARNING:

- You must use a CADD® Extension Set with Anti-Siphon Valve or a CADD® Administration Set with either an integral or add on Anti-Siphon Valve to protect against unregulated gravity infusion that can result from an improperly attached cassette. Unregulated gravity infusion can result in death or serious injury.
- For detailed instructions and warnings pertaining to the Medication Cassette[™] Reservoir or CADD® Administration Set, please refer to the instructions for use supplied with those products.
- Frozen medication must be thawed at room temperature only. Do not heat the Medication Cassette[™] Reservoir in a microwave oven as this may damage the medication, the Medication Cassette[™] Reservoir, or cause leakage.

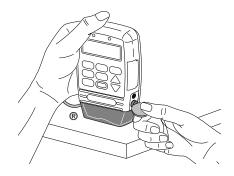
Use aseptic technique as instructed by your clinician.

- Clamp the tubing on the new Medication Cassette™ Reservoir or CADD® Administration Set.
- Insert the cassette hooks into the hinge pins on the pump.



3. Place the pump upright on a firm, flat surface. Press down so the cassette fits tightly against the pump.

4. Insert a coin into the latching button, push in, and turn counterclockwise until the mark on the latch lines up with the solid dot.



WARNING: Attach the cassette (the part of the Medication Cassette[™] Reservoir or CADD® Administration Set that attaches to the pump) properly. An improperly attached or detached cassette could result in unregulated gravity infusion of medication from the fluid container or a reflux of blood, which could result in death or serious injury.

Gently twist and pull on the cassette to make sure it is firmly attached.



- 7. A message will appear on the display showing the type of cassette you have latched. Press (NEXT).
- Reset Reservoir Volume to —
 ? may appear. Press to
 reset the Reservoir Volume
 to the originally programmed
 value. This will also reset the
 Infusion Profile.

High Volume Admin set latched

NEXT to continue

Reset Reservoir Volume to 3000.0 ml?

Press Y or N

Infusion Profile has been reset

Priming the Tubing and Starting the Pump

When the new cassette is attached, the pump *may* automatically prompt you to prime the tubing.

- 1. If Prime Tubing? appears, press .
- This screen will appear. Make sure you are not connected to the pump's tubing. Open the tubing clamps.

Prime Tubing?

Press Y or N

Disconnect tubing from patient Open clamps Hold Y to prime

WARNING: Do not prime the fluid path with the tubing connected to your catheter. This could result in over-delivery of medication or air embolism, which could result in death or serious injury.

 Press and hold the key until the tubing is fully primed or until priming stops.

Priming... 0.1 ml

Hold Y to prime

4. This screen will appear if you release the **\Lambda** key or if priming automatically stops:

Continue Priming?

 If the tubing is not yet fully primed, press the Press Y or N

- **A** key and repeat step 3.
- If you are finished priming, press the W key.

When the new cassette is attached, the pump will automatically prompt you to start the pump.

5. When Start the Pump? appears, check to see if the fluid path is free of air. If your clinician has instructed you to use the Air Detector, go to the next section. If not, follow your clinician's instructions for connecting the tubing to your access site. Open all clamps.

Start the Pump? Press Y or N

6. Press **A** to start the pump.

Starting pump... will appear. The pump will automatically review and display the programmed settings.

RUNNING will appear on the main screen, the green light will blink, and fluid delivery will begin as programmed by your clinician.

* TPN 6230X ***

Starting pump...

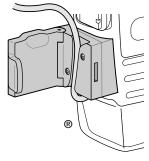
*** TPN 6230X *** RUNNING ↑ Res Vol 1000.0 ml Press NEXT to advance

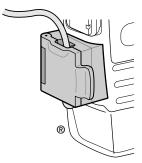
WARNING: Ensure that the entire fluid path is free of all air bubbles before connecting to your catheter to prevent air embolism. Air embolism could result in death or serious injury.

Inserting the Tubing into the Air Detector

The Air Detector is designed to detect air bubbles in the fluid path. When the tubing is inserted into the Air Detector, an air bubble exceeding the specified size will cause an alarm to sound and the pump to stop.

- If your clinician has instructed you to use the Air Detector, open the Air Detector door and thread the tubing through the groove.
- Close the door, making sure the tubing does not get pinched or kinked.
- Follow your clinician's instructions for connecting the tubing to your access site. Open all clamps.





WARNING: Ensure that the entire fluid path is free of all air bubbles before connecting to your catheter to prevent air embolism. Air embolism could result in death or serious injury.

What if I drop or hit the pump?

What should I do if I drop the pump in water? If you accidentally drop the pump in water, retrieve it quickly, dry it off with a towel, and call your clinician.

What if I drop the pump or hit it against a hard surface? Immediately do the following:

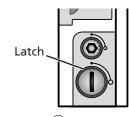
- Check the latch on the side of the pump and make sure the dot on the latch lines up with the solid dot on the pump.
- Gently twist and pull on the cassette to make sure it is still firmly attached.
- Check the battery door to make sure it is still firmly attached.

If the cassette or battery door are loose or damaged, do not use the pump. Immediately stop the pump, close the tubing clamp, and contact your clinician.

WARNING: If the pump is dropped or hit, inspect the pump for damage. Do not use a pump that is damaged, or is not functioning properly. Depending on the type of damage,

death or serious injury could result from the use of a dam-

aged pump.





If there are alarms or special messages you need to be aware of, the pump will beep or sound an alarm. Look at the screen and follow the steps in this table, or press ? for help.

When you see:

Take this action:

9 volt Battery Low

NEXT to continue

The 9 volt battery is low but the pump is operable. Change the 9 volt battery soon.

9 volt Battery Depleted

Install good battery

Install a new 9 volt battery. The pump will not run with a depleted 9 volt battery. A good battery must be installed even when an external source of power is connected.

(blank display or random characters)

A two-tone alarm is sounding and the amber light stays on. The 9 volt battery is depleted. Install a new battery.

Power Pack Depleted

Change Power Source

The Power Pack is depleted. Recharge the Power Pack.

Cassette Damaged Free flow may occur Clamp Tubing NEXT to silence The pump detects the cassette is damaged. Close the tubing clamp and inspect the cassette for damage. Replace it if necessary.

When you see:

Take this action:

Air in line detected

Pump will not run NEXT to silence There is air in the tubing or the tubing is not threaded through the air detector. Press NEXT to stop the alarm. Then follow your clinician's instructions for priming.

Delivery Stopped

(Model 6101 only)

Fluid is not flowing from the fluid container to the pump.
Check for a kink, a closed clamp, or an air bubble in the tubing between the fluid container and pump. Press (STOP) to stop the pump and silence the alarm for 2 minutes, then remove the obstruction and press (NEXT) to restart the pump.

High Pressure

Pump Stopped

There may be a kink in the tubing or a clamp may be closed. Unkink the tubing or open the clamp and the pump will resume delivery. You may press (STOP) to stop the pump and silence the alarm for 2 minutes. After you remove the cause of high pressure, start the pump if necessary. If the alarm continues, contact your clinician.

When you see:

Take this action:

(any screen)

You hear 9 beeps: the infusion profile has ended. Disconnect or change the CADD® Administration set as appropriate.

Reservoir Volume is zero

NEXT to continue

The Reservoir Volume has reached 0.0 ml. Press NEXT to stop the alarm. Then change the fluid container if appropriate.

Reservoir Volume Low

NEXT to continue

The Reservoir Volume value is low, indicating that the level of fluid in the fluid container is low.

Upstream Occlusion

Press STOP to silence

(Model 6101 only)

Fluid is not flowing from the fluid container to the pump.
Check for a kink, a closed clamp, or an air bubble in the tubing between the fluid container and pump. Press (STOP) to stop the pump and silence the alarm for 2 minutes, then remove the obstruction and press (NEXT) to restart the pump.

When you see:

Take this action:

Upstream Occlusion

Press STOP to stop Press NEXT to restart

(Model 6101 only)

Fluid is not flowing from the fluid container to the pump. Check for a kink, a closed clamp, or an air bubble in the tubing between the fluid container and pump. Press (START) to stop the pump and silence the alarm, then remove the obstruction and press (NEXT) to restart the pump.

Error Detected E (error code)

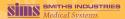
There is a problem with the pump. Close the tubing clamp and remove the pump from service. Contact your clinician.

Your Clinician's
Name:
Phone Number:
Instructions:
Pump placement during bathing/showering:
Pump placement during sleep:
Storage of medication:

Notes	



SIMS Deltec, Inc., St. Paul, MN 55112 U.S.A.



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These products are covered by one or more of the following: U.S. Patent Nos. 4,559,038; 4,565,542; 4,650,469; 5,181,910; 5,338,157; 5,364,242; 5,485,408; 5,531,697; 5,531,698; 5,538,399; 5,540,561; 5,564,915; 5,567,136; 5,567,119; 5,695,473 (Model 6101 only); other patents pending.

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