User Manual

for use with the

PainSmart™ IOD

Ambulatory Infusion System

Moog Medical Devices Group
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Important Safety Note

- When using a Curlin Infusion Pump in a life sustaining situation, use AC Adapter, and provide an extra set of “C” cell batteries in addition to the batteries that are installed in the pump.
- Check pump between patient use per procedure (see maintenance chapter).
- Prior to Programming Load Dose, refer to Important Information in the appropriate Therapy chapters.

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PainSmart™ IOD
Ambulatory Infusion System

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TABLE OF CONTENTS

INTRODUCTION 1

Delivery Mode and Routes 1

Indications for Use 1

Special Features of the *PainSmart™ IOD* Pump 2

Regulatory Agency Approvals 3

Warnings, Cautions, Notices 4

List of Abbreviations 8

FEATURES OF THE PUMP AND ADMINISTRATION SETS 11

Pump Illustrations and Features 11

Disposable Administration Set Illustrations and Features 13

Preparing Medication for Infusion/Changing IV Bags 14

Opening and Closing the Pump Door 16

Choosing and Installing the Curlin Medical Administration Set 17

Gravity Priming Set and Filter 17

Steps in Installing Administration Set into Curlin Infusion Pumps 18

Up Stream Occlusion Alarm Feature 19

Installing the Batteries and Using External Power 20

Curlin Pump Battery Replacement 22

External Power Sources 25

Battery Pack 25

AC Adapter/Charger 31

Bolus Cord Use 32

LEARNING PUMP FEATURES 33

Keypad Function 34

Description of the Info-On-Demand (IOD) Feature of the Moog Curlin Infusion PainSmart™ IOD Pump 35

Info-On-Demand (IOD) Key Functions 36

Description of the Custom Features 39

Improved Pump Display 40

LED Light Indicators 41

Audio Indicator 41

Display Screen 41

BASICS OF PROGRAMMING 43

Types of Display Screens 43
Sample of an Action Field Screen ... 43
Sample of a Selection List Field 43
Sample of a Data Entry Screen 44
Sample of a Run Screen 44
Sample of User Notification Screen 44

Using "Help" 45

Starting the Pump 45
Welcome Screen 45
Power Status Screen 46
Preventive Maintenance Screen 46
Select Program or Bio-Med Setup Screen 46

Access Code 47
Clinician Access Code Screen 47

BioMed Setup 47
Bio Med Setup Menu 47
Enable Therapy Menu 49

Patient History Log 49
History Event Log 50

TABLE OF CONTENTS

Clearing the History Log 51

Clearing Patient Information..... 51

Changing Date and Time 52
Change Date and Time Screen 52

Program 53
Program/BioMed Setup Menu.....53
Resume/Repeat Rx/New Program Menu 53

Resuming a Therapy 54

Repeating a Pre-Programmed Therapy 54

How to Go to "New Program" 55

Entering or Changing Data Fields..... 55

Programming PainSmart pump with Protocol Library Safety System 55

PCA IV THERAPY..... 60

Prescription Menus..... 60
PCA-IV Pre-Prescription Menu Screen 60
Administration Routes.....61
PCA medLIMITS™ Setup Screen..62
PCA Prescription Menu.....63
Run Options Screen.....64

Prime..... 64
Prime Direction Screen.....65

Options 60
PCA Options Menu 60
Clinician Dose Menu.....61
Clinician Dose Run Screen.....61
Shift Totals Screens.....61
Hourly Totals Screen.....62
Clear Hourly Totals Screen 69
PCA Titrate medLIMITS™ Screen 69
Air-In-Line Sensor Off Message Screen 64
Pump Will Sense 2ml Alert Screen 70

Reviewing or Changing the Program 71

Starting the Infusion.....71
PCA Run Screen.....71
PCA Information Screens 72

Interrupting an Infusion..... 72
Pause Menu Screen 73

Resuming the Current Container.. 73
Run Options Screen.....73

Hanging a New Container 73

Entering a New Program 74

Infusion Complete..... 75
PCA Infusion Complete.....75

Stopping the Infusion..... 75
Pause Menu Screen 75

PCA EPIDURAL THERAPY.. 72

Prescription Menus 72
EPI Pre-Prescription Menu Screen 72
Administration Routes.....73
EPI medLIMITS Setup Screen 79
EPI Prescription Menu.....80
Run Options Screen.....81

Prime 81
Prime Direction Screen 82

Options 83
EPI Options Menu.....83
Clinician Dose Menu 84
Clinician Dose Run Screen.....84
Shift Totals Screens.....84
Hourly Totals Screen 80
Clear Hourly Totals Screen 80
EPI Titrate MedLIMITS™ Screen 81
Air-In-Line Sensor Off Message Screen 82
Pump Will Sense 2ml Alert Screen 82

Reviewing or Changing the Program 88

TABLE OF CONTENTS

Starting the Infusion 88
EPI Run Screen 88
EPI Information Screens.....89

Interrupting an Infusion 85
Pause Menu Screen.....90

Resuming the Current Container .. 90
Run Options Screen 85

Hanging a New Container 90

Entering a New Program..... 91

Infusion Complete 92
PCA Infusion Complete 92

Stopping the Infusion 92
Pause Menu Screen.....92

TROUBLESHOOTING 90

Alerts, Alarms, and Error Codes 90

ACCESSORIES 114

Detachable Pole Clamps.....114

Remote Bolus Cord (350-2026) 115

Lockable Safety Shells 116

Pump Holster (340-0111).....117

Keypad Covers 118

CLEANING, ENVIRONMENTAL CONDITIONS, AND MAINTENANCE..... 119

Cleaning.....119

Transport and Storage 120

Operating Environment.....120

Maintenance 120

Limited Warranty.....120

Procedure for Checking Pumps Between Patients 122

TECHNICAL SPECIFICATIONS.....126

CLINICIAN INFORMATION AND ACCESS CODE 134
Clinician Access Code.....134

Lock Setting Table 135

PAINSMART™ IOD CATALOG NUMBERS.....136

TABLE OF FIGURES

Figure 2.1 Curlin Infusion Ambulatory Pump – Front, Top and Left View 11

Figure 2.2 Curlin Infusion Ambulatory Pump – Back, Bottom, and Right View12

Figure 2.3 Sample of Curlin Infusion Administration Sets.....13

Figure 2.4 Spiking the Fluid Container, Gravity Priming the Set and Using the Slide Clamp to Close the Tubing.....14

Figure 2.5 Curlin Infusion Administration Set "Integral Flow-Stop" with Breakaway Tab 15

Figure 2.6 Intentionally Opening the "Integral Flow-Stop" 15

Figure 2.7 Opening the Pump Door 16

Figure 2.8 Closing the Pump Door 16

Figure 2.9 Proper Installation of the Curlin Infusion Administration Set into the Pump.....18

Figure 2.10 Top Cutaway View of the Curlin Infusion Pump with Curlin Infusion Administration Set in Proper Installation Position.....19

Figure 2.11 Curlin Pump Battery Replacement 21

Figure 2.15 Attaching and Removing the Remote Bolus Cord..... 32

Figure 3.1 Curlin Pump Features 33

Figure 3.2 6000 CMS PainSmart™ IOD Keypad with Info-On-Demand Feature 33

Figure 3.3 Sample of Display Screen..... 42

Figure 4.1 History Event Log..... 50

Figure 5.1 PCA Administration Route Default Settings.....57

Figure 6.1 PCA Administration Route Default Settings.....74

Figure 8.1 Detachable Pole Clamp 109

Figure 8.2 Remote Bolus Cord 110

Figure 8.3 Safety Shells 111

Figure 8.4 Pole Clamp and Safety Shell Installation.....112

Figure 8.5 Pump "Holster".....112

Figure 8.6 Keypad Covers 113

Introduction

Let us first review the delivery modes and routes, indications for use, special features, regulatory agency approvals and warnings, cautions and notices relating to the ***Curlin*** *Infusion Pumps.*

Let's begin by introducing you to the **Curlin Infusion** pump, an easy-to-use, ambulatory or pole mounted infusion device designed to meet the needs of the hospital or alternate site patient. It operates accurately in any position, has a low rate resolution of 2 microliters, and its small, compact, lightweight design allows mobility for ambulatory patients.

The **Curlin Infusion** pump can be carried in any of the four convenient carryall soft-packs, pole mounted, placed into the locked "safety shells," or set on a suitable surface for use.

Delivery Mode and Routes

PainSmart™ IOD is a **PCA or Patient Controlled Analgesia** pump designed for therapies that require a continuous rate of infusion, patient-controlled demand boluses, or both. Three Administration Routes, namely intravenous (IV), epidural (EPI) and subcutaneous (SQ); and three Units of delivery, namely milligrams (mg), micrograms (mcg) and milliliters (ml) are available. With the help of a software program purchased from **Moog**, an institution can configure the Administration Route to be fixed as IV, EPI or SQ. If any Administration Route is selected as "Fixed," it is the only Route available to the user (other Routes will not be displayed) and cannot be re-enabled in the Biomed Setup menu. Similarly, the Units of delivery can be fixed as mg, mcg or ml. If any Unit of delivery is selected as "Fixed," it cannot be modified by the clinician/user in the Biomed Setup menu.

Indications for Use

The **Curlin Infusion** pump can be used for intravenous, epidural, or subcutaneous therapies. If Epidural is selected, the Therapy Identifier bar in the Prescription Menu will display EPI instead of PCA. It can be used to deliver medications from the specially designed **Curlin Infusion** medication reservoir, syringes[1] or from IV bags. A physician or a certified, licensed, healthcare practitioner must oversee any therapy. Patients and caregivers using the **Curlin Infusion** pump should be instructed in its use by a qualified clinician and demonstrate an adequate level of proficiency in the use of the pump.

[1] Some syringes are not qualified for use. Please call Moog for details.

Special Features of the *PainSmart™ IOD* Pump

- Small, accurate, ambulatory, volumetric infusion pump that provides reliable and safe delivery of infusion therapies.
- The Info-On-Demand feature of the Curlin Infusion PainSmart™ IOD pump safely provides additional information to the user when the pump is in the "RUN" mode.
- Customized Maximum Basal Rate Limit.
- Customized Maximum Bolus and Dose Limits.
- Customized Lock Level setting.
- Large screen with Tallman Lettering.
- Larger font, single information display.
- Patient side (downstream) "Line Pressure" displayed.
- medLIMITS™ features.
- PCA dose and medLIMITS™ schedules that can be maintained on Repeat.
- Pump will not permit the user to enter any programming value that is outside of its predetermined default range, or further restricted through medLIMITS™.
- Ability, at the institution level, to configure the pump for specific Administration Route or Units of Measure:
    - Units (Factory, ml, mg, mcg, ml fixed, mg fixed or mcg fixed);
    - Route (Intravenous, Epidural, Subcutaneous, Intravenous fixed, Epidural fixed or Subcutaneous fixed).
- Patient and therapy specific programming features via the **Curlin Infusion** BIOMED SETUP and Options Menus (when not limited by the institution).
- Selection by user from three units of delivery, milliliters, milligrams, and micrograms (when not limited by the institution).
- Choice of five or nine digits Clinician Access Code.
- Maintenance Due Date that is automatically set, the first time, after 12 hours of use.
- Shift Totals that can be greater than 24 hours.
- Hourly Totals.
- Automatic scrolling of the programmed prescription with "quick-repeat" function.
- Cost effective, safe and easy-to-load disposable **Curlin Infusion** administration sets featuring a unique, spring-activated, self-clamping, set-based "Integral Flow-Stop" device that automatically clamps the tubing when the door of the pump is opened and prevents inadvertent fluid flow.
- User-friendly, easy-to-teach programming that shortens staff inservice time and patient teaching time.
- "Helpful" help screens and display messages.
- Three tamper-resistant lock settings to assist in maintaining patient compliance and safety.
- Retention of programmed infusion settings until cleared by the clinician, thus eliminating the

need to reprogram before each use.

- Mandatory requirement of two independent key press actions before a running pump can be turned off, thus minimizing the accidental interruption of a therapy in progress.
- Powered by two readily available, cost-effective "C" size alkaline batteries.

IMPORTANT INFORMATION

- Attractive yet robust design that is impact resistant and water resistant.
- Audio alarms that can be adjusted from 1 (very quiet) to 9 (loudest) to meet specific patient and clinical setting needs but that cannot be totally disabled.
- Four sizes of convenient carry packs to assist the ambulatory patient in maintaining independence of lifestyle.
- Retained memory features that allow the pump to easily resume a therapy exactly from where it left off when it is interrupted before completion or to repeat a therapy using the same programmed prescription.
- Compact, lockable safety shells that provide additional security when needed and that are designed to be free standing, pole mounted, or placed in the convenient, soft, carry case.
- Backlit illumination for display screen. Backlight remains on continuously if AC Adapter is in use.
- Integral Up as well as Down Occlusion alarms to notify user if an occlusion is present on either side of the pumping chamber.
- Integral High Upstream pressure alarm to notify user of excessive upstream pressure.
- Integral Air-In-Line alarm to notify user of presence of air in the tubing.
- Adjustable Down Occlusion settings of "High" and "Low" to customize the pump's down pressure requirements.

Regulatory Agency Approvals

INFUSION PUMP
WITH RESPECT TO ELECTRIC SHOCK, FIRE
AND MECHANICAL HAZARDS ONLY
IN ACCORDANCE WITH UL60601-1/CAN/CSA C22.2 NO. 601.1
70TK

- Underwriters Laboratories has evaluated the **Curlin Infusion** Pump with International AC Adapters (360-2022KIT) to UL 60601-1 and CSA C22.2 No. 601-1 for General Standards of Safety for Medical Equipment and found the **Curlin Infusion** Pump to be compliant.
- Underwriters Laboratories has evaluated the Curlin Infusion Pumps with accessory equipments Battery Pack (**350-9262**) and the International AC Adapters (360-2022KIT)to IEC 60601-2-24. Particular requirements for safety of infusion pumps and controllers and they were found to be compliant.
- Underwriters Laboratories has evaluated the **Curlin Infusion** Pump and the AC Adapter (360-2022KIT) for Electromagnetic Compatibility, to regulations EN 60601-1-1-2/1993 and

EN 60601-2-24/1994 and both were found to be compliant.

The indicator shown at the left appears throughout this manual to emphasize important information in the operation of the Curlin Infusion Pump. Please read these sections carefully.

Warnings, Cautions, Notices

EXPLANATION FOR SYMBOLS on the **Curlin Infusion** Pump with the Classified UL 60601-1 and CSA C22.2 No. 601-1 Label.

The meanings for the symbols on the pump are as follows:

⚠ Attention, consult ACCOMPANYING DOCUMENTS.

Type CF Equipment

IPX1 Protected against dripping water.

INFUSION PUMP
WITH RESPECT TO ELECTRIC SHOCK, FIRE
AND MECHANICAL HAZARDS ONLY
IN ACCORDANCE WITH UL60601-1/CAN/CSA C22.2 NO. 601.1
70TK

INTERNALLY POWERED OR FOR USE WITH AC ADAPTER
340-2022/360-2022 and 340-2026/360-2026

DEFIBRILLIATION PROOF TYPE CF APPLIED PARTS

EQUIPMENT PROTECTED AGAINST DRIPPING WATER, IPX1

EQUIPMENT NOT SUITABLE FOR USE IN THE PRESENCE OF A FLAMMABLE ANESTHETIC MIXTURE WITH AIR OR WITH OXYGEN OR NITROUS OXIDE

MODE OF OPERATION CONTINUOUS

WARNINGS

- Prevent direct exposure to radiations. Secure an essential infusion pump outside of the radiation field using extension tubing if required.
- Use only **Curlin Infusion** administration sets in your **Curlin Infusion** pump. Use of nonproprietary administration sets may alter accuracy in delivery of fluids and could result in over or under infusions, leading to possible patient injury or death.
- Visually inspect the pump, pumping chamber and administration set before use. Do not use any pump or administration set that appears to be damaged or tampered with or if there is any indication of improper function.
- Do not use this pump with a pressure cuff applied to the IV bag or medication reservoir bag, and do not unduly squeeze or compress the bag during a running infusion.
- Remove all air from the administration set and IV bag or reservoir before connecting it to a patient's access site.

- Damage to the pump due to dropping or impact while the door is open can cause intermittent free flow conditions and loss of pump accuracy, but may not be visible through inspection. Volumetric testing is the only method to confirm proper function.

  NOTE: Anti-siphon Valve (ASV) administration sets can be used to provide additional free-flow protection.

- Do NOT prime the administration set while it is connected to the patient's access device. Doing so could result in overdosing the patient and could cause injury or death.
- Do not attempt to open the pump's housing. All service problems should be referred to an appropriate service technician.
- **Danger:** Risk of explosion if used in the presence of flammable anesthetic or explosive gasses.

CAUTIONS

- U.S. federal law restricts this device to sale by or on the order of a physician or other licensed practitioner.
- Keep the pumping surfaces clean, dry, and free of fluid spillage at all times.
- This pump is fluid resistant and can withstand fluid spillage. It is **not**, however, designed for total submersion as moisture buildup within the case could cause damage to the operating components. Do not use the pump in the shower, sauna, or steam bath, and do not position the pump where it could accidentally be dropped into a container of fluid (e.g., basin, tub, or toilet). Avoid fluid contact with the power port and data port of the pump.
- Do not try to insert foreign objects into any of the pump connectors, as such objects may damage the pump.
- Use only **Curlin Infusion** external power sources to power the **Curlin Infusion** pump. Using other power sources may result in circuitry or microprocessor damage.
- When using the AC Adapter, connect the plug only into a *grounded* AC outlet.
- Use only non-rigid, non-vented IV fluid containers unless an air vent adapter is in place and the container is suspended from an IV pole.
- Do not unduly stretch the tubing of the administration set or leave the tubing in the pump for more than twenty-four hours when the pump is not running.
- Use **Curlin Infusion** administration sets with appropriate air-in-line filters any time the Air-In-Line Sensor of the pump is disabled.
- If any signs or symptoms of infiltration or inflammation are noted at the infusion site, stop the infusion and report it to the appropriate healthcare provider.
- Always use the slide clamp on the administration set as an additional precaution to occlude the tubing before opening the door of the pump.

NOTICES

- The **Curlin Infusion** pump is not intended for the administration of blood or cellular blood products.
- Dispose of all used administration sets in accordance with all applicable regulatory and institutional policies.
- Teach users to check all tubing sites for proper aseptic connections and to check the administration sets for air leaks before and during the infusion.
- To minimize the potential hazard of air bubble formation or "out-gassing," administer all medications at the proper temperatures, and remove medications from refrigerators as specified by the healthcare provider or pharmaceutical manufacturer.
- Follow local governing ordinances and recycling plans regarding disposal or recycling of device components. When the "C" size batteries are to be discarded, do not incinerate them. Dispose of all used administration sets in accordance with all applicable regulatory and institutional policies and directions.
- Adhere to any warnings, precautions, or recommendations stated by drug manufacturers regarding the use of infusion pumps and disposable administration sets in the administration of their specific products.
- Administer all drugs selected for epidural administration in accordance with the indications included in the manufacturer's package insert accompanying the drugs.
- Note that the administration of medications into the epidural space is limited to delivery via specially designed indwelling catheters inserted by a qualified physician. Any patients receiving epidural infusions should be managed and monitored by medical professionals familiar with epidural administrations and their clinical management. **Moog** offers administration sets specifically designed for epidural infusions with 0.22 micron filters, no injection ports, and yellow striping to clearly identify the set for epidural use. Any administration set being used for epidural infusions should be clearly labeled as such.
- Clean the pump, remote bolus cord, AC adapter, and battery pack with a soft, clean cloth dampened in any of the following:

  1 Warm soapy water (do not submerse)
  2 Isopropyl alcohol
  3 Household bleach, diluted 9:1 with water
  4 Commercial disinfectant

  When clean, dry pump and other items with a soft, clean, dry cloth.
- The soft carry packs are intended for single patient use. They may be cleaned in the gentle cycle of a washing machine using cold water, no chlorine bleach, and line dried. (They cannot be dried in a clothes dryer.)

- The Bolus/Data connector is to be used by the patient only for the connection of the bolus cord with remote switch. Other accessory equipment connected to this digital interface must be certified to the respective IEC/EN standards (i.e. IEC 950 for data processing equipment and EN 60601-1 for medical equipment.) Furthermore, all configurations shall comply with the system standard EN 60601-1. Anyone who connects additional equipment to the signal input part or signal output part configures a medical system, and is therefore, responsible that the system complies with the requirements of the system standard EN 60601-1. If in doubt, consult the technical services department or your local **Moog** representative.
- The **Curlin Infusion Pump** has been tested and found to comply with the European Standard EN 60601-1-2:1993 and EN 60601-2-24:1994. These limits are designed to provide reasonable protection against harmful interference in a typical medical installation.
- This equipment generates, uses, and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to other devices in the vicinity. However, there is no guarantee that interference will not occur in a particular installation. If this equipment does cause harmful interference to other devices, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one of more of the following measures:
  1 Reorient or relocate the receiving device.
  2 Increase the separation between the equipment.
  3 Connect the equipment into an outlet on a circuit different from that to which the other device(s) are connected.
  4 Consult the manufacturer or field service technician for help.

Note: For information regarding the clinician access code, read the chapter entitled, "Clinician Information and Access Code." When entered correctly, the access code allows a user to change the lock settings of the pump and thereby change the settings of the pump.

IMPORTANT INFORMATION

Because the chapter entitled, "Clinician Information and Access Code" contains access code information, the entire chapter should be removed before initial patient teaching and should not be left in the manual while in the patient setting. Doing so could jeopardize the security levels intended only for clinician use and access.

List of Abbreviations

#BOLS/hr = Number of Bolus Doses per hour
In Patient controlled analgesia the number of boluses allowed per hour 0-15

Δ RATE = Amount rate may be changed

Δ TIME = Interval until rate may be changed

AMT = Amount

AMT = Amount to be infused
TBI

BA = Bolus attempted
Options hourly screen shows amount of bolus doses attempted

BASL = Basal
Basal rate or continuous basal infusion maximum varies by route of infusion, intravenous, epidural or subcutaneous

BG = Bolus given
Options hourly screen shows amount of bolus doses infused

BOL INT = Bolus Interval
Allows clinician to enter RX amount of time 0-60 min between patient-activated bolus dosing for pain management

CLIN = Clinician
Clinician dose is the option to give an additional prescribed dose by a clinician

CLRHRLY = Clear hourly
Option to clear hourly totals of amount infused

CONCEN = Concentration
Concentration is the amount of milligrams or micrograms per milliliter of solution

dd = Day
To set date enter 2 digit day

DN Occlu = Down Occlusion
Occlusion pressure in the line between the pump and the patient access site, as a kink in tubing, blockage in access site (infiltration) or clamp on

EPI = Epidural
To inject anesthetic into the spinal canal

HH:MM = Hours and Minutes
To set time enter Military 24 hour or AM/PM 12 hour

HX = History
The patient history log is a cumulative record of the last 6,000 events that have occurred in the pump

INF = Infusion
Fluid or medication pumping into the patient via intravenous, subcutaneous, or epidural routes

IV = Intravenous
To give fluids or medication directly into the venous circulation via peripheral or central circulation

KVO = Keep vein open
To infuse a very small amount of fluid to keep vein patent

LPRES = Line Pressure
In CMS-enabled pumps shows downstream pressure to patient in millimeter of mercury

MAX = Maximum

MCG = Microgram

medLMTS = Sets soft limits for medication being infused

MG = Milligram

MIN = Minimum
Minimum amount of time between patient boluses

ML = Milliliter

A unit of liquid measure 1/1000 of a liter

mm = Month
To set date enter 2 digit month

MX = Maximum
Upper limit of a bolus or patient demand dose, or infusion rate

OPT = Options
Once prescription is programmed, this key provides customizing features as locking programs from changes, alarm volume, battery level, amount of air detection, adjust occlusion alarm, hourly totals, shift totals, clinician dosing

PCA = Patient controlled analgesia
Pain control by continuous rate of infusion with options for patient controlled demand boluses

PT = Patient
Person receiving infusion

RX = Prescription
Written order for medication with instructions of how to administer as time, dose, amount, limits, and route

SENS = Sensor
Air in line sensor will detect the amount of air in the tubing

SQ = Subcutaneous
To give fluids or medication just below the skin, also (SC)

VOL = Volume
Amount of fluid in intravenous bag, medication reservoir or syringe

yy = Year
To set date enter 2 digit year

Chapter 2

Features of the Pump and Administration Sets

*The **Curlin Infusion** PainSmart™ IOD Pumps have been designed with the user in mind and have the latest in friendly, simple-to-learn technology to allow fast and easy access to the features of these pumps. With a little introduction and training, an operator will quickly be able to program and use these pumps to deliver the therapies prescribed.*

Pump Illustrations and Features

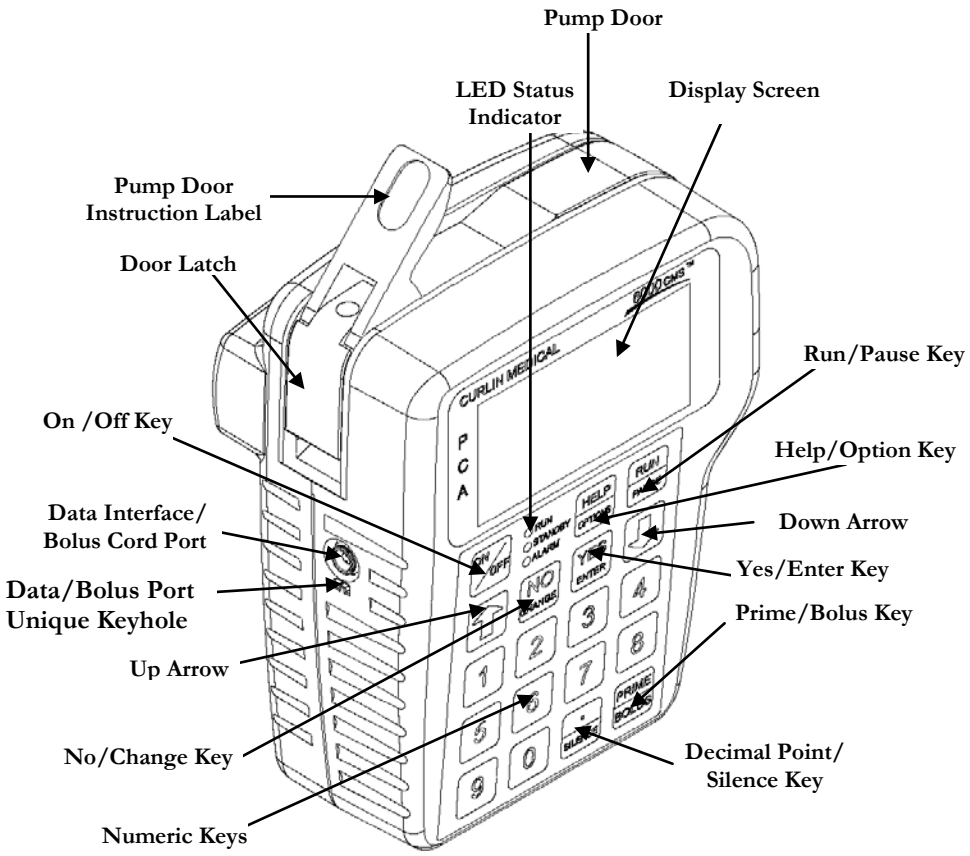


Figure 2.1 Curlin Infusion Ambulatory Pump – Front, Top, and Left View

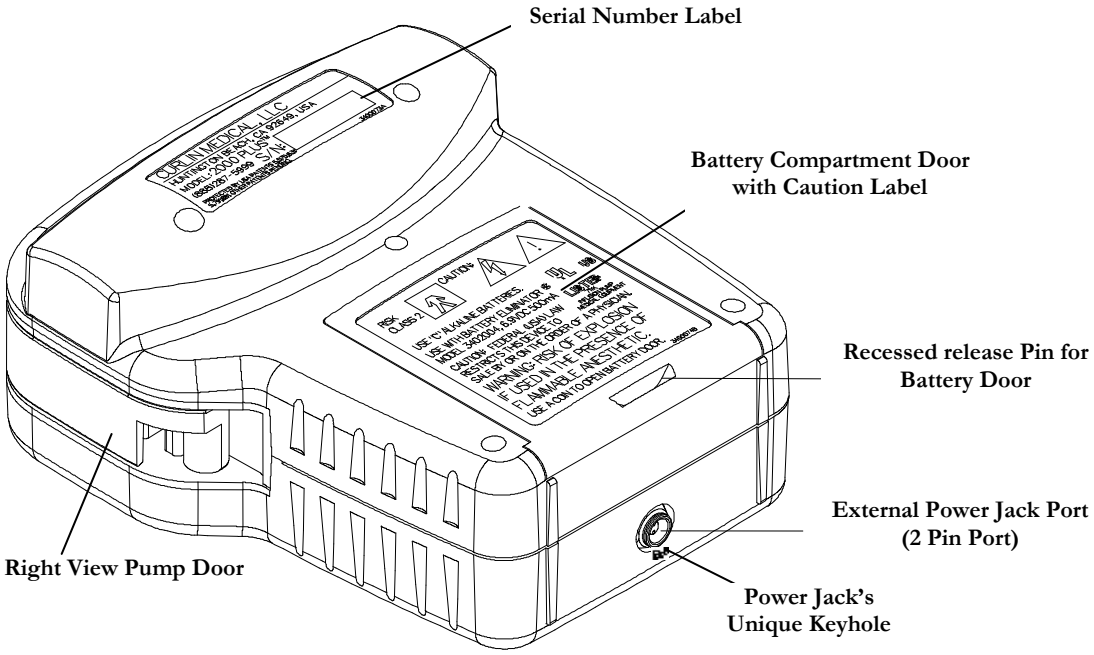
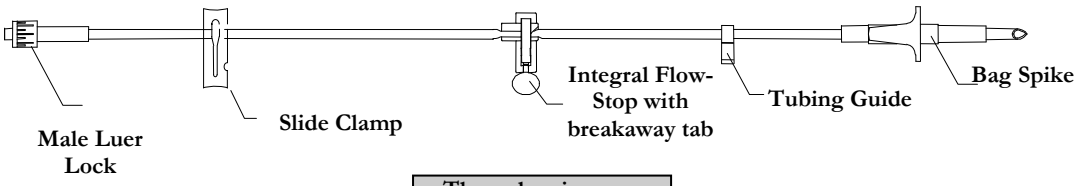


Figure 2.2 Curlin Infusion Ambulatory Pump – Back, Bottom, and Right View

Disposable Administration Set Illustrations and Features



These drawings are NOT to scale.

Note: An important feature of the Curlin Medical administration set is the unique design of the set-based "Integral Flow-Stop." When the set is new, the "Integral Flow-Stop" has a breakaway tab that keeps the "Integral Flow-Stop" open and allows the tubing to be gravity primed. When you are ready to load the administration set into the pump, remove this tab, and the "Integral Flow-Stop" will automatically clamp the tubing. The Curlin Medical "Integral Flow-Stop" prevents inadvertent flow of medication to the patient whenever the door of the pump is opened because the "Integral Flow-Stop" automatically reacts to clamp the tubing. The "Integral Flow-Stop" can, however, be opened after the tab is broken away by intentionally squeezing down on the movable spring action section of the "Integral Flow-Stop." See Figure 2.6 for an illustration of how to intentionally open the "Integral Flow-Stop." For a complete listing of all available administration sets, see the Curlin Medical Product Catalog.
Note: Administration sets with anti-siphon valve protection can also be used for additional free-flow protection.

CAUTION: USE ONLY CURLIN INFUSION ADMINISTRATION SETS WITH THIS PUMP

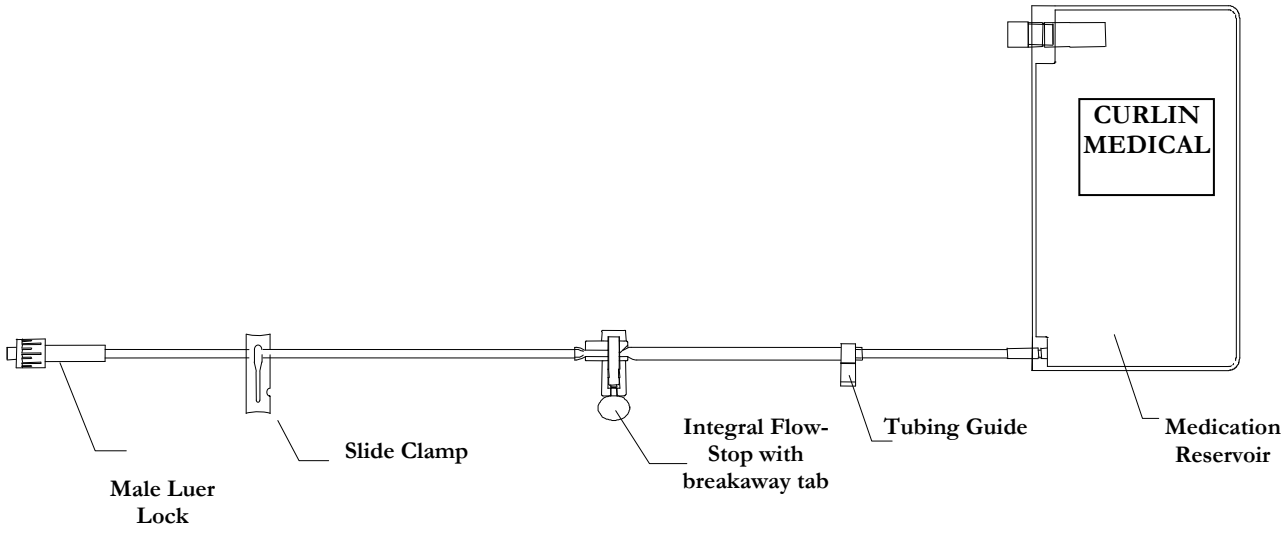
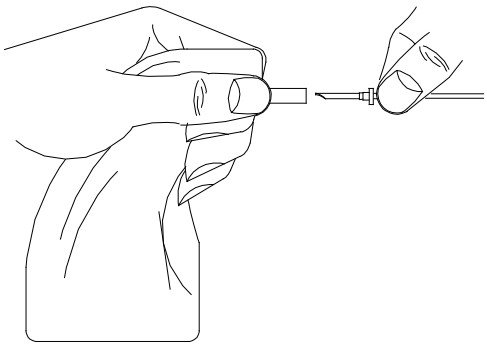
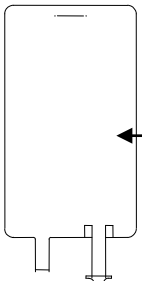


Figure 2.3 Sample of Curlin Infusion Administration Sets

Preparing Medication for Infusion/Changing IV Bags

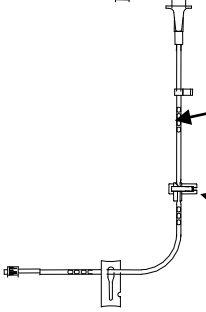


When using an IV bag, grasp the bag spike firmly in one hand and the IV bag spike-port in the other, and insert the spike completely into the port using aseptic technique.



Holding the bag with fill ports down allows fluid to fill tubing and displace the air in the line. Be sure all air has been removed from the IV bag as well.

Air in the line moves down through the tubing as the fluid fills.



Notice the breakaway tab on the "Integral Flow-Stop" has been removed. The set can be gravity primed by intentionally squeezing the "Integral Flow-Stop" and thus releasing the clamping action. (See Figures 2.5 & 2.6)

OPEN CLOSED

In these illustrations, the slide clamps are shown in their open and closed states.

Figure 2.4 Spiking the Fluid Container, Gravity Priming the Set, and Using the Slide Clamp to Close the Tubing

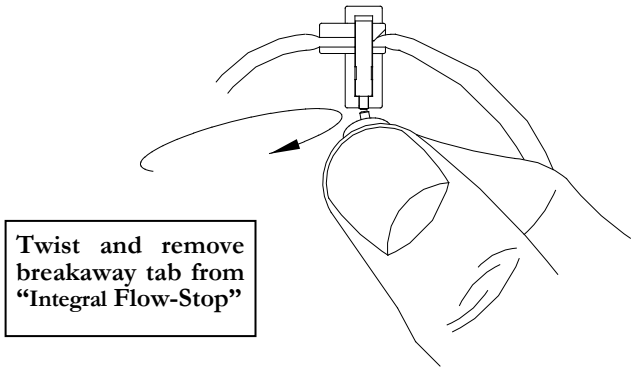


Figure 2.5 Curlin Infusion Administration Set "Integral Flow-Stop" with Breakaway Tab

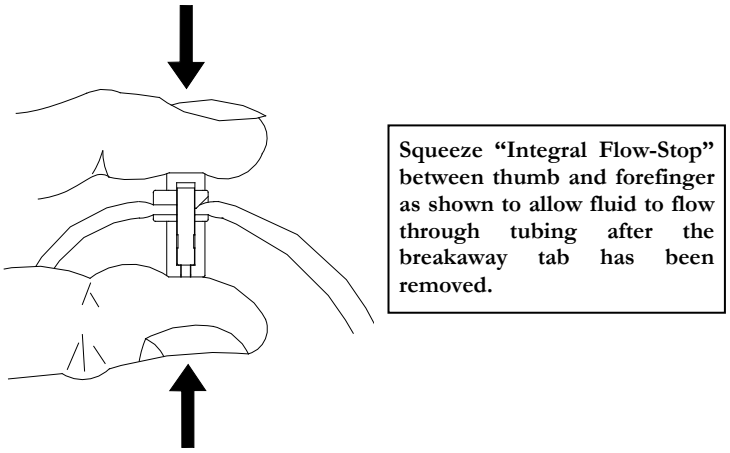
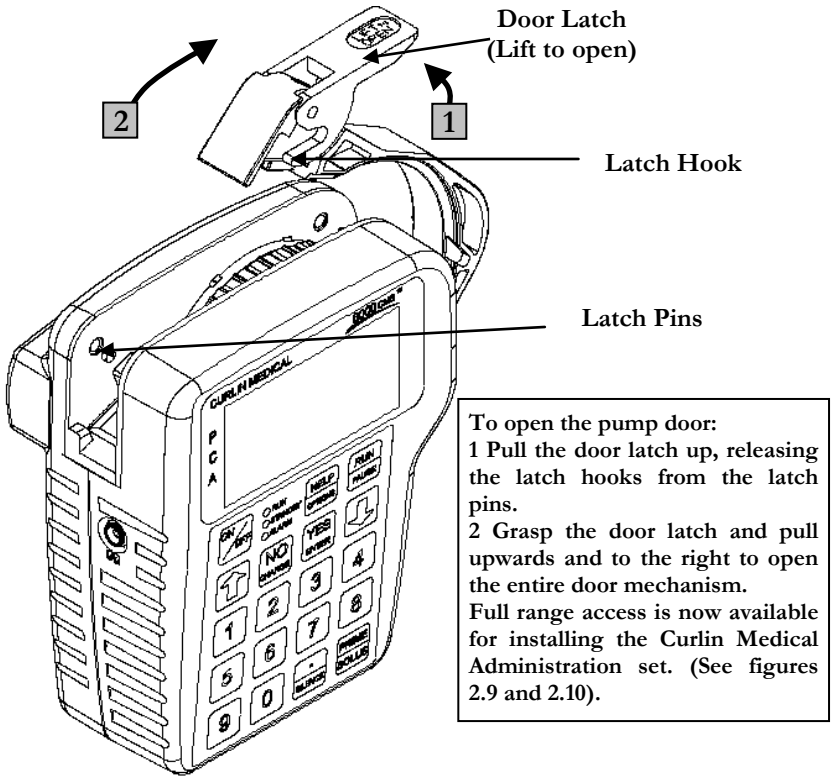


Figure 2.6 Intentionally Opening the "Integral Flow-Stop"

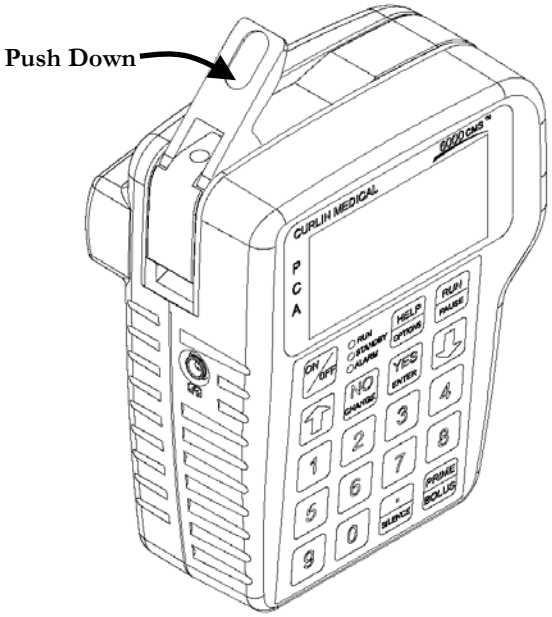
Opening and Closing the Pump Door

The pump door, located on the top of the **Curlin Infusion** pump, has a sturdy "over center" latching mechanism, which assists in securely closing the door. Review the following illustrations:



To open the pump door:
1 Pull the door latch up, releasing the latch hooks from the latch pins.
2 Grasp the door latch and pull upwards and to the right to open the entire door mechanism.
Full range access is now available for installing the Curlin Medical Administration set. (See figures 2.9 and 2.10).

Figure 2.7 Opening the Pump Door



To close the pump door properly, move the entire door assembly to the left and push downward until it is in its closed position. Then push the door latch down until it locks. The "over-center" latching mechanism assures that the door closes easily and securely.

Figure 2.8 Closing the Pump Door

Choosing and Installing the Curlin Infusion Administration Set

A number of distinctively designed **Curlin Infusion** administration sets are available to deliver each prescription accurately. (See **Curlin Infusion "*Product Catalog.*"**) Each set is designed to meet the requirements of specific therapies, therefore, if a basic bag-spike set or a set with a medication reservoir is needed, with or without air eliminating filters, etc., there are several configurations from which to choose. Follow the healthcare provider's protocols for preparing the medication and the administration set before loading the set into the pump, remembering the following basic rules:

1 Examine the packaging and the administration set before use to assure the package integrity. Do not use the set if the protective caps are dislodged or if the package integrity is breached, as fluid path sterility cannot be guaranteed under those circumstances.

2 Notice that there is an instructional insert included in the administration set package. This insert includes written and graphic instructions for proper installation of the administration set into the pump. It also lists the "fill volume" of the administration set. Allow for the amount of the fill volume when determining the final IV bag volume.

Gravity Priming Set and Filter

1 To gravity prime the administration set, twist and remove the breakaway tab from the "Integral Flow-Stop", then spike the IV bag. Squeeze the "Integral Flow-Stop" between your thumb and forefinger to allow fluid to flow through the tubing until all air is removed from the IV bag and administration set. (See figures 2.5 and 2.6.) Release the "Integral Flow-Stop" to automatically clamp the tubing. **To correctly prime a filtered set, hold the filter in a vertical position with the flow downwards allowing the filter to completely fill with fluid.**

Note: Administration sets with integral anti-siphon valves cannot be gravity primed.

2 Close the slide clamp on the administration set before the door of the pump is opened. This step along with the automatic clamping properties of the "Integral Flow-Stop" prevents inadvertent fluid flow.

3 Use an administration set with an air-eliminating filter whenever the Air-In-Line Sensor is disabled or set to "OFF."

4 Follow CDC guidelines or Intravenous Nursing Standards of Practice Guidelines regarding the frequency of changing the administration sets.

5 Use aseptic technique and universal precautions as directed by the policies of your healthcare facility or institution.

6 Use only **Curlin Infusion** administration sets with the **Curlin Infusion** pump.

7 Dispose of administration sets in accordance with all applicable regulatory and institutional policies and directions.

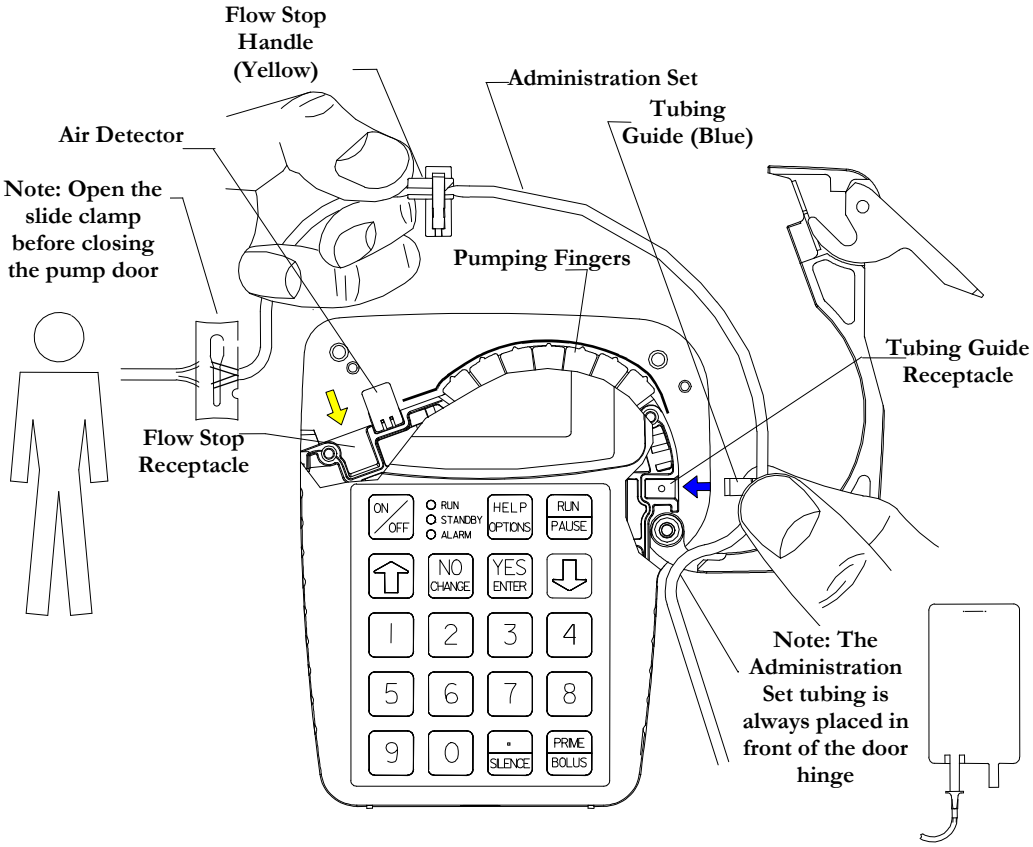
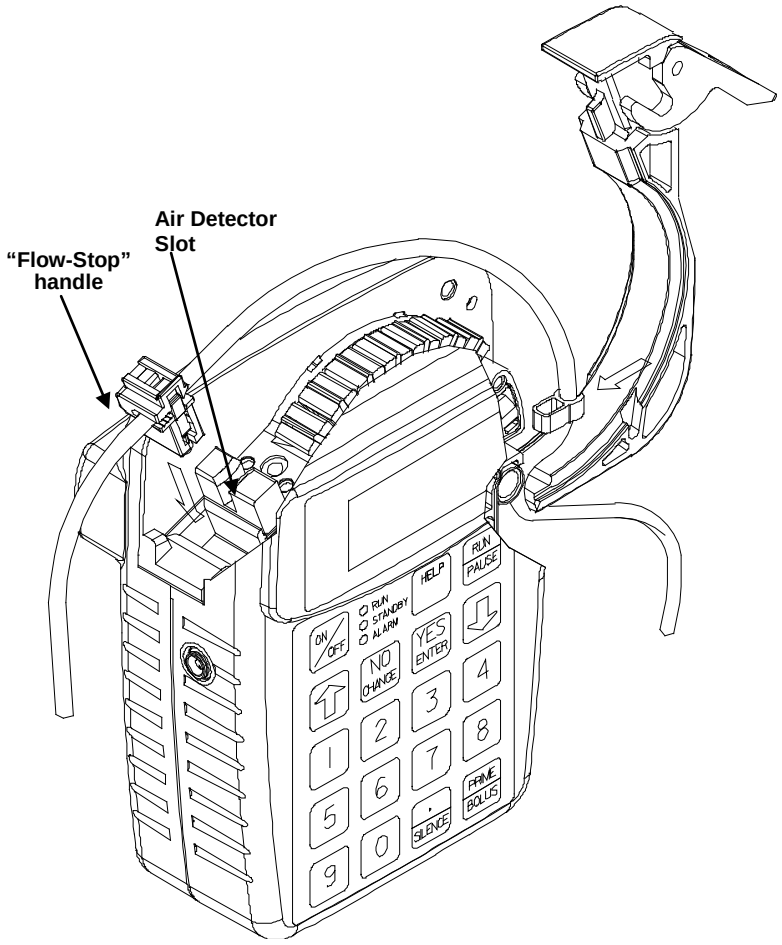


Figure 1.9 shows the Curlin Medical pump with a cut-away of the display screen allowing you to visualize the receptacles for the small *blue* "Tubing Guide" on the right side of the pump and the larger *yellow* "Integral Flow-Stop" on the left side of the pump. The arrows, in corresponding colors, are visible on the background surface of the pump head and indicate the proper direction for placing these locators into their receptacles. Also, note that the breakaway tab has been removed from the "Integral Flow-Stop", allowing the administration set to be placed into the pump correctly.

Figure 2.9 Proper Installation of the Curlin Infusion Administration Set into the Pump

Steps in Installing Administration Set into Curlin Infusion Pumps

1 Fully open the door of the pump.
2 Close the slide-clamp on the administration set.
3 Activate the Integral Flow-Stop by twisting and removing the breakaway tab.
4 Insert the blue tubing guide into the receptacle on right side of pump in the direction of the blue arrow, positioning the tubing in **front** of the door hinge.
5 **Center the tubing in the middle of the pumping fingers.**
6 Holding the yellow Integral Flow-Stop by the handle (see Figure 2.9), insert it on an angle into the receptacle on the left side of the pump in the direction of the yellow arrow. **CAUTION: Do not press down on the top of the Integral Flow-Stop when inserting it into the receptacle. This action could inadvertently open the Integral Flow-Stop and allow fluid to flow through the tubing.**
7 Push the tubing slightly into the black Air Detector slot.
8 Open the slide-clamp on the administration set.
9 Fully close the door of the pump and latch it securely.
10 If the set is not primed, prime the set according to prime section in the relevant therapy chapter.
11 Attach the primed set to the patient's venous access device per agency protocols.
12 Dispose of used administration sets in accordance with all applicable regulatory and institutional policies and directions.



1 Insert blue "Tubing Guide" on right side of pump with tubing in front of the door. The Blue arrow on the background surface of the pump head indicates direction and location for the blue Tubing Guide.

2 Important: Center the tubing horizontally over the pumping fingers covering the yellow dot.

3 Insert yellow "Integral Flow-Stop" on the left side of the pump. The yellow arrow on the background surface of the pump head indicates the direction and location for the "Integral Flow-Stop."

CAUTION: Hold the "Flow-Stop by the small handle. DO NOT press down on the top of the "Integral Flow-Stop" when inserting it into the receptacle. This action could inadvertently open the "Integral Flow-Stop" and allow fluid to flow through the tubing.

4 Press the tubing into the air detector slot gently with index finger.

Figure 2.10 Top Cutaway View of the Curlin Infusion Pump with Curlin Infusion Administration Set in Proper Installation Position

Up Stream Occlusion Alarm Feature

The **Curlin Infusion** pump has an Up Stream Occlusion alarm feature, e.g., it detects low up-stream pressure. This feature notifies you if an occlusion is present between the IV container and the pump. If this alarm occurs, the display will read, "ALARM –UP OCCLUSION," the alarm will sound, the red LED light will blink, and the infusion will stop. This situation requires that you press the "Pause" key, examine the administration set from the IV container to the pump for potential causes of the occlusion. If a clamp is in place, open, or remove it; if the administration set is kinked, straighten it. If indicated, close the slide clamp on the administration set and open the door of the pump to verify that the set is correctly placed. Close the pump's door and reopen the slide clamp. When ready to begin infusing, select "RESUME" and then press the "RUN" key.

The pump also detects excessive up stream pressure (pressure on the IV bag). The display will read "ALARM – HIGH UP PRESSURE," the alarm will sound, the red LED light will blink, and the infusion will stop. If this situation occurs, relieve the source of excessive pressure on the IV bag and when ready to begin infusing, select "RESUME" and then press the "RUN" key.

Note: The up stream occlusion detection is disabled during priming.

Installing the Batteries and Using External Power

The **Curlin Infusion** pump can be powered solely with the use of two "C" size alkaline batteries installed into the pump as shown in Figure 2.11. "C" size batteries are readily available at most retail facilities that carry alkaline batteries. A second alternative for power is the portable, rechargeable Battery Pack.. The third alternative for power is to use the AC Adapter, allowing the pump to be connected to any approved, three-prong electrical wall outlet.

The "C" size batteries will last approximately 30 hours at a rate of 125 ml/hour and over 85 hours at rates of 2 ml/hr or less. Factors that influence battery life are the number of times the screen is illuminated with a key press, the viscosity of the fluid being delivered, the age and temperature at which the batteries have been stored. To extend the life of the batteries, patients can use the AC adapter whenever they are stationary, e.g., at night.

The pump has a "low battery" alert that beeps periodically with a message appearing to tell you when the power is low in the "C" size batteries. This alert gives you a window of time to be sure you have back-up power (e.g. extra batteries, AC adapter, or a recharged battery pack.) At a continuous infusion rate of 25 ml per hour, you can expect at least a 30 minute warning before your pump can no longer operate due to empty batteries. At higher infusion rates your advance warning will be less.

The user should be taught that whenever the "low battery" alert appears, to pause the therapy, turn the pump off, and to replace both "C" size batteries. Then, turn the pump back on, select resume to continue the therapy.

When the "empty battery" alarm occurs the pump will go into an alarm state and the infusion will stop. If this occurs, turn the pump off, replace both of the "C" size batteries, turn the pump on, and go to the "Resume" function to continue the therapy. All programming information is retained. If the "C" size batteries are empty and you want to connect the AC adapter to power the pump, turn the pump off first, plug in the AC adapter and then turn the pump back on.

Note: If you are operating the pump solely on the battery pack or the AC Adapter (no "C" size batteries installed) and the power source is disconnected from the pump, the alarm will sound, the infusion will stop, and the display screen will be blank. Turn the pump off, reconnect the power source, turn the pump on, and resume the therapy. All programmed information is retained. However, as there is no low battery alarm for the external battery pack, the external battery pack should always be used with charged "C" cell batteries should batteries.

When replacing batteries, **pause any infusion in progress by pressing the "PAUSE" key, and then turn the pump off.** Replace the batteries, and, when the pump is turned back on, select "RESUME" to continue the therapy from the point at which it was interrupted.

Figure 2.11 Curlin Pump Battery Replacement

STEP 1 - The battery door compartment is located on the back of the pump.

To open the door, move the dark gray slider located at the bottom of the door to the right "Unlock" position until you hear a click.

STEP 2 - Slide the battery door down and remove it from the pump.

STEP 3 - If your pump contains a plastic tab between the two batteries, pull up on this tab until batteries can be removed. Otherwise follow the instructions below.

If it does NOT contain a plastic tab between the batteries, use a penny or dime to pry the top (+) edge of each battery out.

Do not use a screwdriver or other sharp object to pry batteries out of the pump.

To check battery levels while the pump is running:

- Press the Help/Options keys
- Scroll down to PWR Ck?
- Press no to change to ON.
- Press yes to view battery level
- Press Yes to Accept OPT? to ext the Options menu

Battery Replacement, Alternate Battery Door

- The battery door compartment is located on the back of the pump.
- To open the door, depress the recessed release pin. (Use a small coin such as a penny to depress the pin.)
- **Do not use a screwdriver or other sharp objects to open battery compartment door.**

- Slide the battery door down and remove it from the pump.

- If your pump contains a plastic tab between the batteries, pull up on the tab until the batteries can be removed. Otherwise:
- Using the small coin (Penny) pry up and out on the top (+) edge of the battery to cause the battery to pop out.
- Use the same technique on the second battery.
- **Do not use a screwdriver or other sharp objects to pry batteries out of the pump.**

- Install two new "C" size batteries by depressing the coil spring with bottom (–) end of the battery and sliding the top of the battery into place.
- Please note: Both positive poles are at the top of the battery compartment. To depress the coil spring, insert the battery at a slight angle.

- Slide the battery door compartment back to its closed position until the recessed pin locks into place.

To check battery levels while the pump is running;**

- Press the Help/Options keys
- Scroll down to PWR Ck?
- Press no to change to ON.
- Press yes to view battery level
- Press Yes to Accept OPT? to exit options

CAUTION: If batteries are removed from the bottom (–) end first, damage to the springs in the bottom of the battery case can occur. Damaged springs can create a situation in which the battery overheats. If the battery or the

menu	pump becomes excessively warm to the touch, do not attempt to remove the batteries. Set the pump aside and contact your home infusion pharmacy.

* It is recommended to use Duracell® Procell®[1] Batteries in Curlin Pumps
** "C" cell batteries only, this function will not check external battery pump power levels.

[1] Duracell and Procell are registered trademarks of Proctor & Gamble.

External Power Sources

Rechargeable Lithium Ion (Li-Ion) Battery Pack (360-2030)

Overview

The Li-Ion Battery Pack (pictured below) is designed to be used with 2 fully charged C-cell batteries to power a model 2000, 4000, 6000 or PainSmart™ infusion pump for an extended period of time without being connected to a power outlet. The battery pack allows the user to operate the pump for a minimum of 24 hours (at 125ml/hr) in addition to the run-time supplied by the disposable C cells in the pump. To ensure an adequate low battery warning period, the battery pack should always be used with 2 fully charged C-cell batteries installed in the pump.

The battery pack provides 5.2V with a capacity of 6000 mA-hrs. It is connected to the same port connector that the AC adaptor uses to power the pump. When the battery pack power is fully exhausted the pump will automatically revert to the internal C batteries. Having new C cell batteries in the pump is crucial to ensure an adequate low battery warning period and to ensure infusion continuity if the battery pack is accidently disconnected. The Red Low Battery LED on the battery pack will illuminate prior to the the external battery running out of power. The pump will provide at least 30 minutes of audible and visual warning prior to the internal C cell batteries running out power.

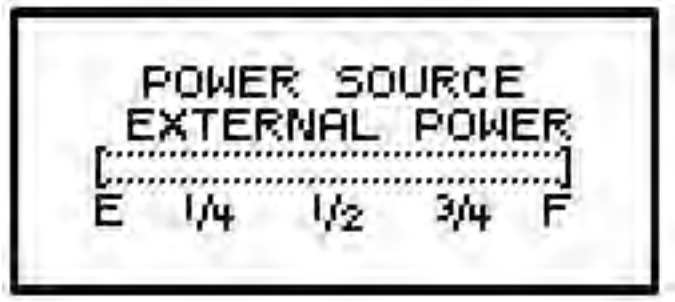
The battery pack is recharged by connecting it to an AC adaptor. Fully recharging the battery takes approximately 15 hours. While charging, the LED indicator will be yellow. When the battery is fully charged, the LED will be green. If a fault is encountered in the battery pack the LED will either flash yellow or not illuminate.

Warnings

- The external battery pack should be used with new C cell batteries in the pump to avoid interruption of the infusion if the cord between the pump and battery pack becomes disconnected and to ensure proper operation of the pump's low battery alarm.
- Do not use this battery pack with the POWDEC AC Adaptor WM10-6.9V (Curlin p/n 340-2022) as the combination may exceed the allowable limits for emitted electromagnetic interference.
- Battery pack cable utilizes an attached cable. This can be a tripping or chocking hazard, avoid use that may lead to entanglement around the neck or feet.

Cautions

- This battery pack is fluid resistant, but it is not designed to be water tight. Care should be taken to avoid any unnecessary contact with water or other fluids. Do not use the battery pack where it might fall into water.
- The power gauge located on the pump screen (pictured below) does not reflect the remaining charge in the battery pack. The displayed capacity on the pump will always show empty when a battery pack (with remaining capacity) is attached (see image below):



- Refer to the LED indicators on the pump to determine if the battery pack has been fully charged or is in a low battery condition.
- The battery pack should be unplugged from the pump when recharging. The AC adaptor cannot simultaneously power the pump and charge the battery pack. If AC operation is desired, connect the adaptor directly to the pump.

Notice

The Rechargeable Battery Pack DFU is a supplement to the Moog Medical Devices Group Pump User's Manual. All other applicable WARNINGS, CAUTIONS and NOTICES are found in the User's Manual and should be referred to when using this DFU.

Environmental

Storage Temperature -20ºC to 65ºC, Atmospheric pressure 500hPa to 1060hPa, Humidity 10% to 90%
Operational Temperature 5ºC to 40ºC, Atmospheric pressure 650hPa to 1020hPa, Humidity 10% to 90%

Directions for Use

Charging the Battery Pack

The Curlin 360-2022 (Globtek GTM41076-0609-1.5), Curlin 340-2022 (Globtek GTM41076-0609-1.5), Curlin 360-2026 (Globtek GTM341-7.5-800) or Curlin 340-2026 (Globtek GTM341-7.5-800). AC adapters can be used to recharge the battery. A fully discharged battery will recharge in less than 15 hours.

- If connected, disconnect the battery pack from the pump.
- Connect the AC adapter to the power port of the battery pack.

- The charge LED will illuminate yellow if the battery needs to be recharged. See below:

- Charge the battery pack for 15 hours or until the LED turns Green indicating the battery is fully charged.

- Disconnect the battery from the AC adaptor.

  Note: If there is a fault detected while charging the battery, the LED will either flash yellow or not illuminate. Most faults are due to either the AC adapter becoming unplugged from the power outlet or the battery pack being at a location where the temperature is not between 5 and 40 degrees Centigrade. If the fault condition persists, then the battery pack should be returned to Moog Medical Devices Group for replacement of the battery.

Using the Battery Pack to Power the Pump

- Make sure fully charged C cell batteries are loaded in the pump.
- Connect the battery pack to the AC adapter port on the pump.

- Operate the pump in the desired infusion mode.

- Do not use the pump's display (see cautions) to predict external battery life.
- When the external battery pack is nearly depleted, the red LED indicator will be illuminated.

- The pump may continue to be used until the disposable C cells are depleted.
- Disconnect battery pack from pump prior to recharging the battery pack.

Maintenance

- There is no need to fully discharge the batteries periodically since Li-Ion batteries do not have a memory effect. The battery pack enclosure may be cleaned with a mild soap or disinfectant.

End of Life Determination

The battery in the battery pack degrades with charge cycles and time since manufacture. When the batteries are no longer holding their charge for a sufficient length of time, the battery pack should be replaced.

AC Adapter/Charger

This accessory becomes the external power source when plugged directly into the **Curlin Infusion** pump. The AC Adapter can also be used to recharge the portable battery pack by plugging the connecting cable into the recharging port on the battery pack.

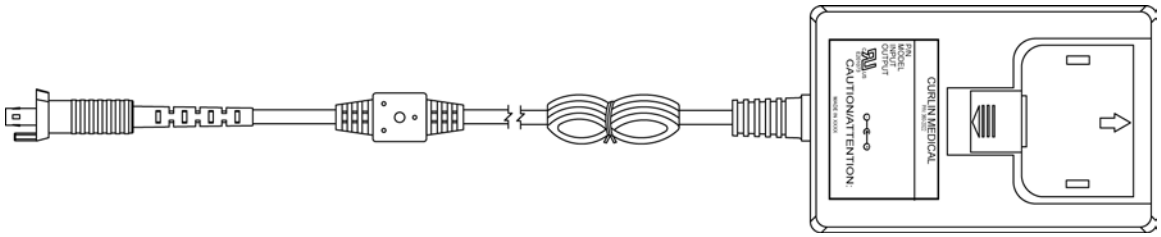
There are two AC Adapters:

- The US Domestic AC Adapter (P/N: 360-2026)has a three-prong plug to be plugged into a standard three-prong, grounded wall receptacle and a six-foot cable which plugs into the power outlet on the bottom of the pump. The two-pin connector plugs into the receptacle by aligning the red dots on each.
- The International AC Adapter (P/N: 360-2022Kit) available for international use. The two-pin connector plugs into the receptacle by aligning the key on the connector.

Figure 2.13 AC Adapters

Note: Pause the pump and then turn it off before connecting the AC Adapter.

Note: The connector is keyed with a "notch" at this red-dot location to guide alignment. To remove, pull back on the locking sleeve to unlock and then pull gently to remove the connector from the battery pack or pump's port.



International AC Adapter

Figure 2.13 Universal AC Adapter

Two-Pin Adapter plugs into port on bottom of the pump for AC power or can be used to recharge the battery pack by plugging into the recharging port on the battery pack. To remove, pull back on the locking sleeve.

Bolus Cord Use

When the PCA therapy is used, patient bolus dosing can be programmed. To facilitate ease of patient self-dosing, the patient can use the Bolus key or the Remote Bolus Access Cord. When a bolus dose is requested one beep will sound to indicate that the dose has started. If a bolus dose is requested and denied, two beeps will sound. When a bolus dose is completed, no beep will occur.

The cord is five feet long and has a patient access button at the top of the remote cord. The four-pin adapter with the unique key plugs into the four-hole port with the matching keyhole on the left side of the pump (see Figure 2.14 below). This port may be covered with a plastic plug. If so, remove the plug to access the port. To remove the cord, pull back on the soft, green locking sleeve and disengage the cord from the pump by pulling it straight out of the access port.

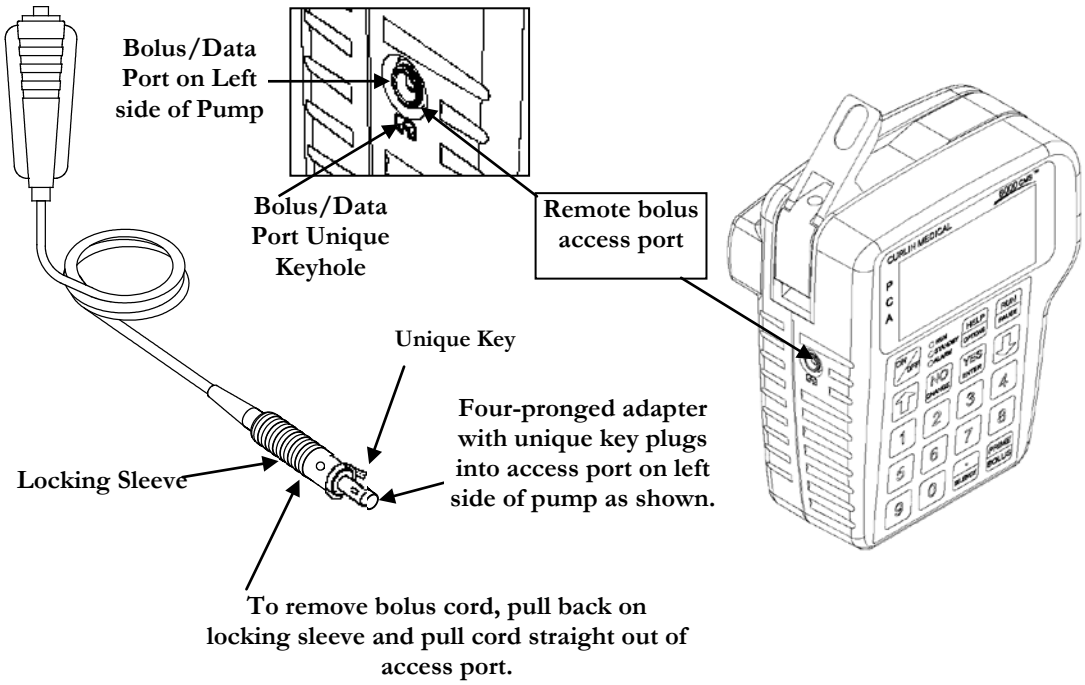


Figure 2.15 Attaching and Removing the Remote Bolus Cord

Chapter

3

Learning Pump Features

Learning to operate the ***Curlin Infusion*** *pump requires a few basic points of understanding that remain constant in every therapy. It may be helpful to read through these steps first, and then go back and use them to operate the* ***Curlin Infusion*** *pump.*

First, examine the key pad and the functions of the individual keys. Knowing how the keys, especially the dual function keys, work will allow you to give commands to the pump.

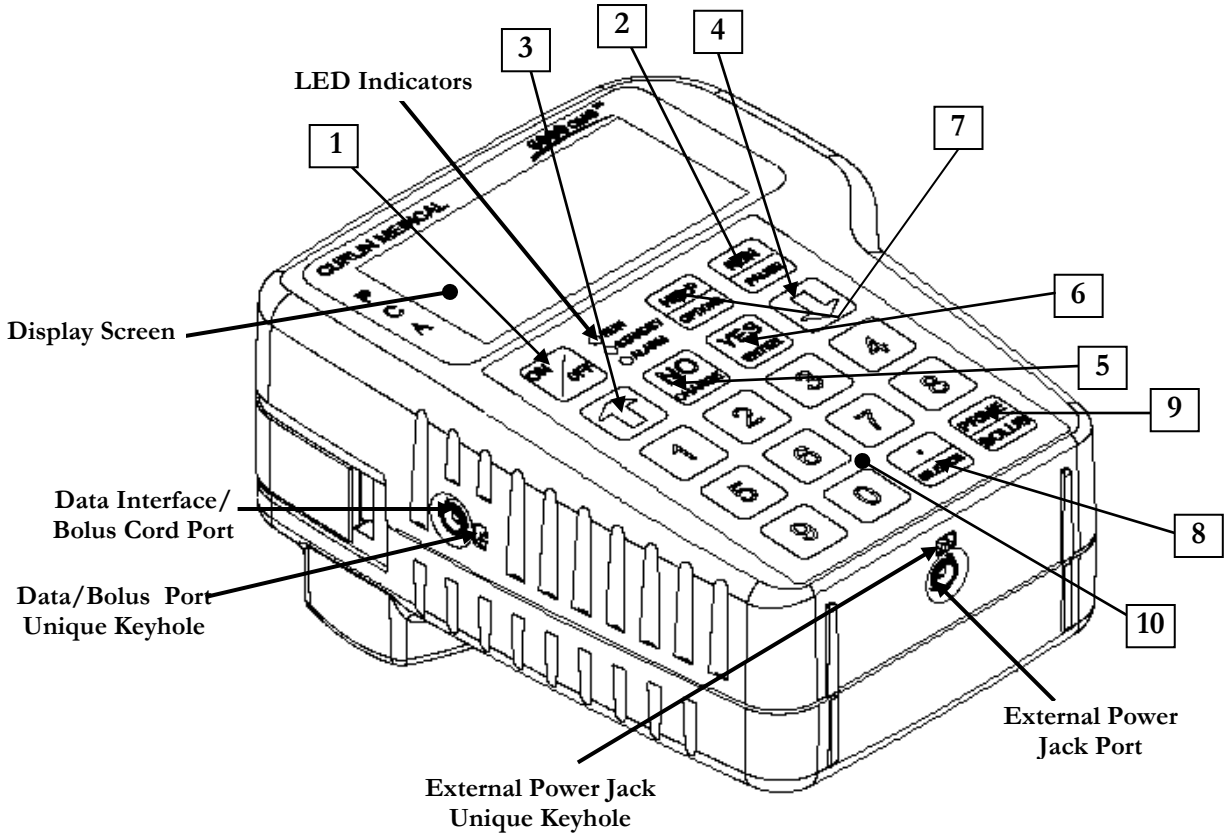


Figure 3.1 Curlin Pump Features

Keypad Function

1 **ON/OFF:** Used to turn the pump on and off. When the pump is running, however, the "PAUSE" key must be used to stop the infusion first and then the "OFF" key can be pressed to turn the pump off. This requirement prevents the interruption of an infusion by a single accidental press of the "OFF" key.

2 **RUN/PAUSE:** Used to start or pause the motor of the pump, which, in turn, starts or pauses fluid delivery. When the pump is running, pressing this key will pause the pump. When the pump is paused, pressing this key during the "Run Options" menu will start the pump. This key is also used to terminate a pump alarm.

Note: If the cause of the alarm is not resolved when the pump is "Resumed," the alarm will recur.

3 ⇑ **Up arrow:** Used to scroll the highlight bar or field cursor *up* on the display screen.

Note: The Up arrow will not accept a selection which must be made with a YES/ENTER or NO/CHANGE key.

4 ⇓ **Down arrow:** Used to scroll the highlight bar or field cursor *down* on the display screen.

Note: The Down arrow will not accept a selection which must be made with a YES/ENTER or NO/CHANGE key.

5 **NO/CHANGE:** Used to respond "NO" to data or questions presented or to reject or to change highlighted data.

Note: The **Curlin Infusion** pump requires you to actively change a selection. The Up arrow or Down arrow will not change a selection.

6 **YES/ENTER:** Used to respond "YES/ENTER" to data or questions presented or to "accept" highlighted data when prompted to press "enter" key.

Note: The **Curlin Infusion** pump requires you to affirmatively accept a selection. The Up arrow or Down arrow will not accept a selection.

7 **HELP/OPTIONS:** Used to request a help screen when the pump is paused or is at the "Infusion Complete Screen." It is also used to access the options screen when the pump is running. When the options screen is displayed, help is again available with this key. When in a help screen, use the arrow keys to scroll through the help message, and then press the help key again to exit help.

8 **./SILENCE:** Used as a decimal point when programming numeric data. Used also to temporarily silence the alarm (for one-minute intervals).

9 **PRIME/BOLUS:** Used to prime fluid through the tubing to clear it of air or to administer a patient demand bolus dose of medication in PCA therapy.

10 **0 - 9 Numeric keys:** - Used for data entry and the display of Info-On-Demand functions when the pump is in the "RUN" mode.

Description of the Info-On-Demand (IOD) Feature of the Curlin Infusion PainSmart™ IOD Pump

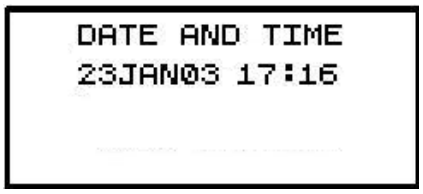
Figure 3.2. PainSmart™ IOD Keypad with Info-On-Demand Feature

To view any of the Info-On-Demand screens on the pump display, press the related key while the pump is in the "RUN" mode, i.e., running a therapy. The information requested will be displayed for 6 seconds and then the display will return to the previous infusion screen. Exception is the "CLRShft /9" key.
The information for the "CLRShft/9" key will be displayed for 11 seconds and it will allow the user to "CLEAR" or "EXIT" within the 11 seconds or the screen will revert to the "RUN" screen automatically.

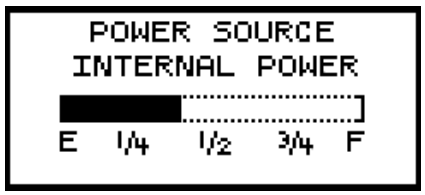
Info-On-Demand (IOD) Key Functions

Designation and functionality of the ten numeric keys of the Info-On-Demand keyboard:

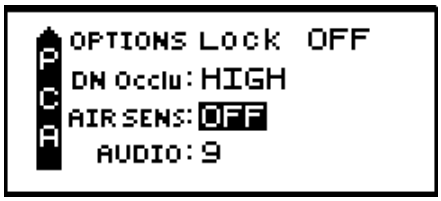
1 **Date and Time (DateTime):** The Date and Time screen displays the current date and time according to the pump. Changes to this screen are not allowed.



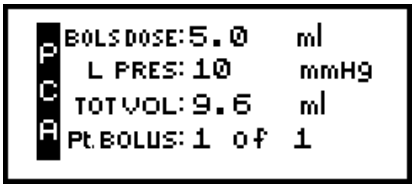
2 **Power Check (PowrChck):** The Power Check screen displays the active power source of the pump, whether internal (batteries) or external. If the active source is the C-size batteries, a bar graph will indicate the amount of power remaining in the batteries. This graph is only an approximation of the charge left in the batteries and not intended to give specific representation of the battery life. Furthermore, the bar graph should NOT be used to check the state of the external NiMH Battery Pack, because it will not provide with accurate results.



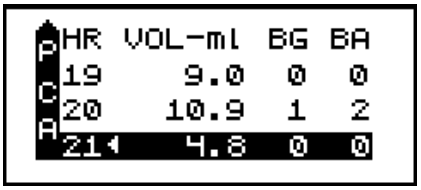
3 **Option Info (OptnInfo):** This screen provides information on selected options. The information displayed here includes the status of the Lock Level, Down Occlusion, Air Sensors, and the Audio Level. Changes to this screen are not allowed.



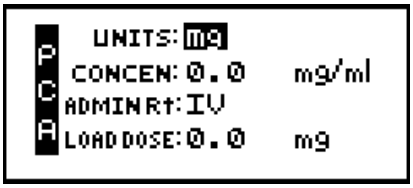
4 **Hold Screen (HoldScrn):** Pressing the Hold Scrn/4 key will hold the current Run screen information for an additional six seconds and then the pump will return to the normal running screen. Changes to this screen are not allowed.



5 **Hourly Total (HrlyTot):** The information displayed here includes the hourly Volume (amount) Infused, the number of Bolus Given and the number of Bolus Asked. Within the allocated 6 seconds, the user can scroll up or down the screen. Changes to this screen are not allowed.

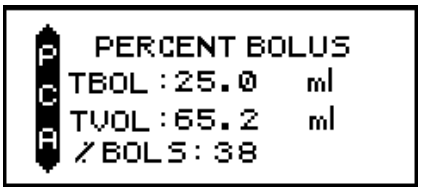


6 **Therapy Information (ThrpyInfo):** The information displayed in this screen is from the pre-prescription menu and includes Units, Concentration, Administration Route and Load Dose. Changes to this screen are not allowed.



7 **Rx Information (RxInfo):** This screen provides prescription menu information such as Basal Rate, Patient Bolus, Bolus Interval, and Number of Bolus per Hour. Changes to this screen are not allowed.

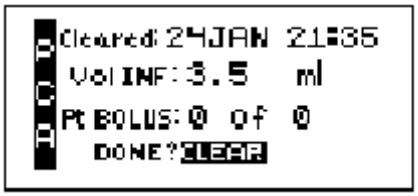
8 **Percent Bolus (%Bols):** This screen provides Bolus Volume as percentage of Total Volume. TBOL is the total volume infused through bolus, TVOL is the total volume infused, and % BOLS is the percentage of TBOL over TVOL. Changes to this screen are not allowed.

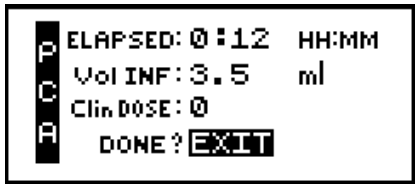


9 **Clear Shift Total (CLRShft):** Two SHIFT TOTAL screens are displayed for 11 seconds when this key is pressed. This key allows the user to view or CLEAR the SHIFT TOTALS (this action does not interfere with the operation of the pump).

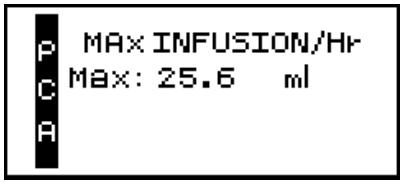
To **CLEAR** the SHIFT TOTAL information to zero, press the CLRShft/9 IOD key, and then press the YES/ENTER button in response to the "DONE? CLEAR" question and the pump will go back to the running screen. Press the CLRShft/9 IOD key again to verify that the values are reset or cleared.

To **EXIT** the SHIFT TOTAL screens without clearing, press NO/CHANGE for the "DONE? CLEAR" question and then YES/ENTER for "DONE? EXIT" and return to the running screen. The pump will return to the running screen automatically after 11 seconds have elapsed since the pressing of the CLRShft/9 IOD key, even if no key is pressed.





10 **Check Maximum Infusion (ChckMax):** The information displayed here is the maximum (rate or amount) that can be infused hypothetically, given the prescription parameters. This "Maximum Infusion per Hour" serves as a "sanity check," after the pump has begun infusing. Changes to this screen are not allowed.



Description of the Custom Features

A customized **Curlin Infusion** PainSmart™ IOD pump may have Maximum Basal Rate Limit and Maximum Bolus and Dose Limits values that are different from previous default maximum values for all administrative routes (IV, EPI, and SQ), and a different Lock Level other than the default value OFF. These values, once programmed at the factory, can not be changed unless the pump is initialized or a new setting is programmed, both of which must be performed by **Curlin** or factory trained biomedical professionals.

1 Customized Maximum Basal Rate Limit

A customized **Curlin Infusion** PainSmart™ IOD pump may come with a Maximum Basal Rate Limit value set in the factory that is different from the default maximum value.

An alert will be displayed if a value greater than the factory set Maximum Basal Rate Limit value is entered in BASL RATE, MedLMTS Mx Rate, and MedLMTS Delta Rate.

For example, while programming a PCA (IV) therapy in a customized **Curlin Infusion** PainSmart™ IOD pump, where the Maximum Basal Rate Limit is factory set to be 9.9 for IV administration route, if you enter 21 for BASL RATE, an alert message will appear:

"GREATER THAN MAX. FIELD LIMIT MAX=9.9"

The values you are allowed to enter in this example, for BASL RATE, are between 0.1 and 9.9, with an increment of 0.1. Same limit will apply to MedLMTS Mx Rate, and MedLMTS Delta Rate also, if the therapy calls for them.

2 Customized Maximum Bolus and Dose Limits

A customized **Curlin Infusion** PainSmart™ IOD pump may come with a Maximum Bolus and Dose Limits value set in the factory.

An alert will be displayed if a value greater than the factory set value is entered in Load Dose, Clinician Dose, Pt. Bolus, and MedLMTS Pt. Bolus.

For example, while programming a PCA (EPI) therapy in a customized **Curlin Infusion** PainSmart™ IOD pump, where the Maximum Bolus Dose Limit is factory set to be 9.9 for EPI administration route, if you enter 10 for LOAD DOSE, an alert message will appear:

"GREATER THAN MAX. FIELD LIMIT MAX=9.9"

The values you are allowed to enter in this example, for LOAD DOSE, are between 0.1 and 9.9, with an increment of 0.1. Same limit will apply to Clinician Dose, Pt. Bolus, and MedLMTS Pt. Bolus also, if the therapy calls for them.

3 Customized Lock Level setting

A customized **Curlin Infusion** PainSmart™ IOD pump may have a certain Lock Level value set in the factory which is not the default, OFF. It can be any one of these values: Lock Level 1, Lock Level 2, or Lock Level 3.

The pump user is not allowed to change the customized factory programmed Lock Level setting. If you attempt to change the customized Lock Level, the pump will display an alert "MODIFICATIONS NOT ALLOWED."

Improved Pump Display

In addition to the above custom features, Tallman Lettering is implemented in this new version of the PainSmart™ IOD pump for better safety.

The PainSmart™ IOD pump displays smaller numerical fonts to the right of the decimal in the PCA therapy programming menus and in the RUN screen menus where the data is displayed. Trailing zeros are dropped, i.e., if a decimal value is 0, then the 0 after the decimal is not displayed. Also, the unit for milliliters is displayed as "mL", instead of "ml." For example, the value ten milliliters is displayed as 10 mL (not 10.0 ml) and the value 10.2 milliliters is displayed as 10.2 mL (not 10.2 ml).

LED Light Indicators

This "three-light" feature gives the user feed back on the current status of the pump.

1 **Green** indicates a RUNNING status of the pump and blinks approximately once every three seconds when a therapy is in progress.

2 **Yellow** indicates a STANDBY status of the pump and blinks while the pump is paused.

3 **Red** indicates a STOPPED or ALARM status of the pump and blinks whenever an alarm status exists.

4 **Green and Red** indicates an "Infusion Complete Alarm" status with KVO or a Down Occlusion Alarm with automatic restart.

5 **Yellow and Green** indicates a "Standby" status with automatic start. This occurs when a "0" basal rate and bolus only is programmed.

Audio Indicator

An audible alarm or "beep" is used to alert you to a change in the status of the pump or a situation requiring user action or notification. A message will appear on the display screen describing the situation. The rate, number of beeps, and volume further specify status. This pump offers nine different volume levels from which to select for the audio feature; however, the "beep" cannot be totally disabled. The audio "beep" can be silenced for one-minute intervals by pressing the "SILENCE" key except during an "Error Code," at which time the alarm continues and the pump must be turned off. The chapter of this manual entitled, *Troubleshooting* includes tables with the alerts, alarms, and error codes, the conditions causing them, the message displayed on the screen, and the actions required for resolution. If an alert or alarm occurs, please refer to these tables.

Display Screen

The display screen provides a maximum of four lines of text, or fields, at any one time for feedback regarding the pump status and data entry. It notifies the user with prompts or messages when necessary. The screen has backlighting that remains:

- on if connected to an AC outlet;
- lit for 15 seconds if using batteries or during any key press sequences,
- on with alert/alarm messages.

When no keys are pressed for 15 seconds, the light goes off to conserve battery power. Backlighting is resumed with any key press.

In most displays, a "Therapy Identifier" Bar appears at the far left of the screen, indicating which therapy is running or which programming screen is being considered. This identifier also has another function, and that is to alert you when more text exists either above the top line or below the bottom line on the display screen. The identifier bar will have an arrow configuration either pointing up or down or both whenever there is more text, or it will appear as a squared-off block if no further text exists.

The screen also has a cursor or highlight bar to tell you which field is being considered. Use the up and down arrow keys to move the cursor. When the pump is running, a specially designed "running screen" appears to provide information regarding the status of that particular infusion. In the following chapters, each display screen will be explained.

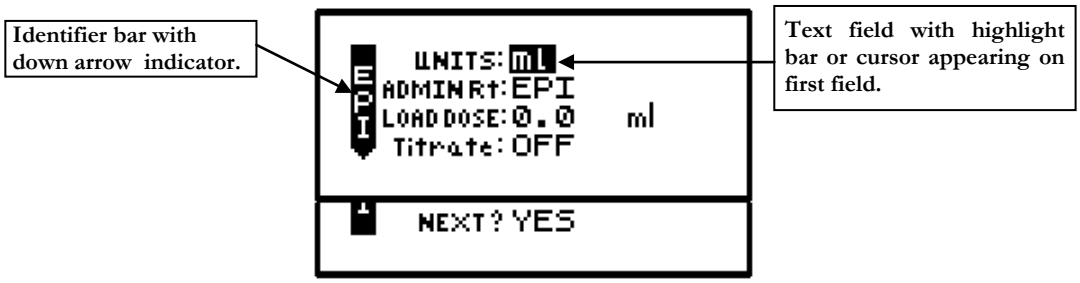


Figure 3.3 Sample of Display Screen

Chapter 4

Basics of Programming

*The **Curlin Infusion** pump allows the operator to customize this device to meet the specific infusion needs of each individual patient. This chapter will explain the customizing features along with the programming steps.*

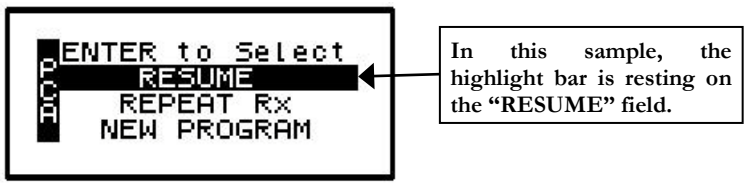
Programming the **Curlin Infusion** pump requires initial orientation and training, but the programming steps have been designed to be user friendly and self-prompting to minimize training time.

Several different types of screens and fields may appear on the displays. Each of the different types of screens is explained below. Whenever you are required to interact with the pump, a highlight bar or cursor will be present on the screen.

Types of Display Screens

1 An "action field screen" allows you to select a pump action from the menu.

Sample of an Action Field Screen



To select a field other than the one highlighted, use the arrow keys to move the cursor up or down to the desired field. When the cursor is highlighting the correct field and you press the "YES/ENTER" key, the pump accepts the selection and executes that action. In the example above, the pump would proceed with steps to resume the therapy.

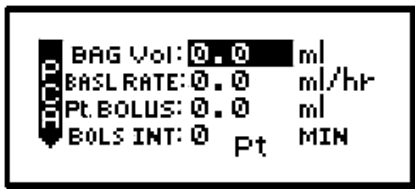
2 A "Selection List" field allows you to select from a predetermined list of values:

Sample of a Selection List Field

The above example allows you to select which unit to use in programming—e.g., Units: ml, mg or mcg. Only one selection is displayed at a time. When the cursor blinks it is indicating that no selection has yet been accepted. The selection displayed on the screen changes each time you press the "NO/CHANGE" key. Pressing the "YES/ENTER" key accepts the selection displayed and moves the cursor to the next field.

3 "Data entry screens" are used to specify prescription parameters.

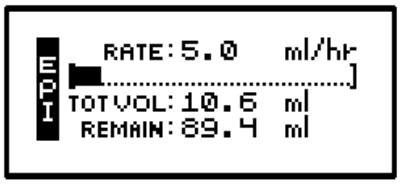
Sample of a Data Entry Screen



In the above example, the cursor rests on a field requiring numeric data entry for specific prescription information, e.g., the volume in the IV bag. The operator enters the correct information using the numeric keys and decimal point key. When the correct data is displayed and the "YES/ENTER" key is pressed, the pump accepts the selection if it is a valid entry value and moves the cursor down to the next field. If the down arrow key is used to move to the next field, the value reverts back to default or the last programmed value.

4 "Run screens" display information to the operator about the progress of the specific infusion while the pump is running:

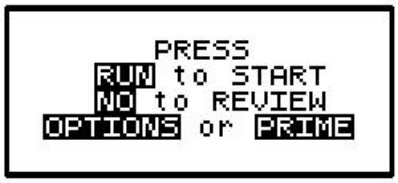
Sample of a Run Screen



The standard "run screen" displays the Mode Identifier, the rate of pumping, a graphical bar that indicates the progress of the infusion, the volume already infused, and the time remaining for the infusion. No action is required on a run screen, therefore, no highlight bar appears. In some therapies a field may toggle to give the operator more information, i.e. the "REMAIN" field may change from *HH:MM* (time remaining) to *ml* to tell you how much volume is remaining as well. The information on the run screen will alternate between a four-line display and a large font display of each line of information on the screen.

5 "User Notification Screens" are messages displayed to the user for information.

Sample of User Notification Screen



Words representing "keys" are displayed in reverse video (shown in this manual as bold text). The action requested by a notification screen is either to press a specified key or to resolve an alert, alarm, or error code.

For all events such as alerts, alarms, or error codes, specific notification screens will also be displayed. (See chapter entitled, *"Troubleshooting"* for specific alert, alarm, and error code messages and how to resolve them). The audio alarm also beeps and the LEDs blink whenever an alarm or error code occurs. In the event of an error code, user directions will be

displayed on the screen, most often, this will require turning the pump off and back on again.

Using "Help"

Help screens are available for data entry/selection screens, during "action field" screens, and at the "infusion complete" screen. To access help, press the "HELP" key. Use the arrow keys to scroll through the help message, and then press the "HELP" key again to exit help. When a help screen is displayed and there is more text to read which is not currently on the screen, a small up (▲) or down (▼) arrow indicator will appear to let you know you can scroll up or down to read the complete help message.

Starting the Pump

Start the pump by pushing the blue "ON" key. The following screen displays:

Welcome Screen

The above "Welcome" screen gives the date, time, and a message that the pump is performing a self-diagnostic test. If the numbers on the date field are displayed as zeros, the system clock will need to be set using the BIOMED Set Up. (See "BIOMED SETUP" section.) The display also lists the software revision number for this pump.

During this test, you will hear three beeps as each of the LED indicators blink. If any one of the green, yellow, or red LED lights fails to light or to blink, return the pump for service. If the self-test fails, the alarm will sound, and a new screen will appear stating: "ERROR CODE:#" or "SYSTEM ERR:#" (See "Troubleshooting" chapter for a list of error notices and how to resolve them). The error notice will include a numeric code. At any error notice, the only user option is to turn the pump OFF.

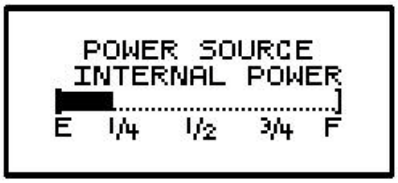
IMPORTANT INFORMATION

If, at any time, there is indication of pump damage, error code, or messages requiring the pump to be turned off, call the healthcare agency or facility as directed to report the condition.

Note: If an error notice appears, turn the pump off and back on again. If the pump passes its start-up diagnostics tests, you can safely resume the therapy. If the error notice recurs, however, turn the pump off and notify the healthcare agency responsible for monitoring the pump. Some error notices may reset the pump back to its default settings and therefore will require reprogramming.

When the **Curlin Infusion** pump successfully passes it's self-test, a brief message of "System OK" appears, and the display moves to the next screen:

Power Status Screen



This screen displays the active power source of the pump, whether internal (batteries) or external power. If the active source is the C-size batteries, a bar graph will indicate the amount of power remaining in the batteries. This graph is only an *approximation* of the charge left and is not intended to give a specific representation of battery life. The display will again automatically move to the next screen:

Preventive Maintenance Screen

The above alert screen will appear after the Welcome Screen ***only*** when the Maintenance Due date has arrived and will continue to appear until the date has been reset during the actual maintenance performance. The pump may continue to be used after the maintenance due date but should be returned for its routine maintenance at the earliest opportunity.

Select Program or Bio-Med Setup Screen

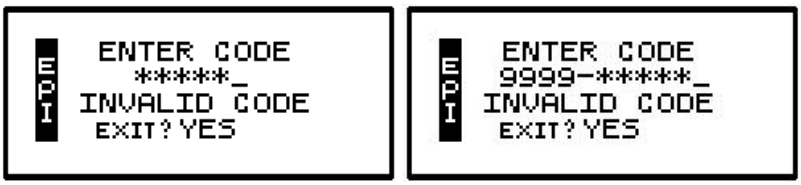
This screen is the first requiring you to act. It gives two selections from which to choose, "PROGRAM" or "BIOMED SETUP." Using the up or down arrows, move the cursor to the desired field and then press the ENTER key.

Selecting *"PROGRAM"* allows you to proceed with choosing features and entering data required for the pump to deliver a prescribed therapy.

Selecting *"BIOMED SETUP"* allows the pump to be programmed and customized by the clinician to perform a number of special functions including setting the system clock. (See "BIOMED SETUP") Accessing "BIOMED SETUP" requires you to enter a clinician access code. (Refer to the chapter entitled, *Clinician Information and Access Code,* of this manual.)

Access Code

Clinician Access Code Screen



Depending on the configuration of your pump, the clinician enters a five-digit access code; or a four-digit personal identification number followed by the five-digit access code; and, if the code is valid, the display will automatically move to the requested screen or field. The four-digit personal identification number appears on the screen but the access code numbers do not; instead five asterisks appear. **If the "four-five" configuration is selected, the personal identification number is recorded in the history file for the purpose of providing an audit trail**. If an invalid code is accidentally entered, a message displays: "Invalid Code" and the clinician can reenter the correct code.

Note: After entering the access code, DO NOT press the "YES/ENTER" key. The pump will automatically recognize the correct code and give you access to the secured section.

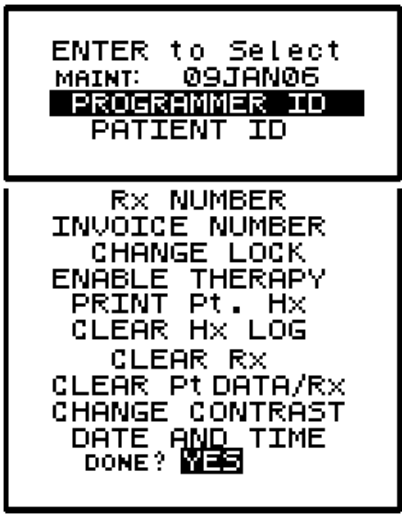
IMPORTANT INFORMATION

If an access code screen displays and <u>the access code is not known</u>, use the down arrow to move the cursor to the "EXIT?" field. Press the "YES/ENTER" key at this field to return to the previous screen and make another selection. Also, remember that the "HELP" key is available when the pump is paused.

Look first at the *BIOMED SETUP* menu and its special customization features:

BioMed Setup

Bio Med Setup Menu



The BIOMED SETUP menu lists special optional customization and performance functions of the pump:

IMPORTANT INFORMATION

You may skip any of these functions by using the down arrow to move the cursor to only the desired selection. When the desired selection is highlighted, press the "YES/ENTER" key. A secondary screen will be displayed. When all BIOMED SETUP functions are completed, move the cursor to the last field, "DONE? YES," and press the "YES/ENTER" key to exit this menu.

1 PREVENTIVE MAINTENANCE DATE: The first field displays the date the pump requires its annual routine preventive maintenance check.

Note: You cannot change this date; it is changed only when a factory technician performs the routine pump maintenance.

The clinician that initially programs the pump for a patient should note the maintenance date. Once this date arrives, a "Perform Routine Maintenance" message will briefly appear after the welcome screen. If the "Maintenance Due" date will occur during the time the patient will be on service, the clinician **may** wish to select another **Curlin Infusion** pump for this patient. This action will avoid any undue concern on the patient's part at the notice of "maintenance due" and eliminate the need to recall a pump in service for maintenance. The pump, however, can continue to be used without disrupting service when this message appears, but should be returned for service as quickly as possible.

2 PROGRAMMER ID: This feature gives the programmer the option to use the numeric keypad to enter his or her assigned ID number (up to ten digits). If this option is not to be used, it can be left blank and skipped, using the down arrow to scroll to the next field. When the PROGRAMMER ID number is satisfactory, press the "YES/ENTER" key at the "ACCEPT? YES/NO" field and the number will be stored in the pump history log.

3 PATIENT ID: This feature gives the programmer the option to use the numeric keypad to enter a patient ID number (up to ten digits). If this option is not to be used, it can be left blank and skipped, using the down arrow to scroll to the next field. When the PATIENT ID number is satisfactory, press the "YES/ENTER" key at the "ACCEPT? YES/NO" field and the number will be stored in the pump history log.

4 Rx NUMBER: This feature gives the programmer the option to use the numeric keypad to enter a prescription number (up to ten digits). If this option is not to be used, it can be left blank and skipped, using the down arrow to scroll to the next field. When the Rx NUMBER is satisfactory, press the "YES/ENTER" key at the "ACCEPT? YES/NO" field and the number will be stored in the pump history log.

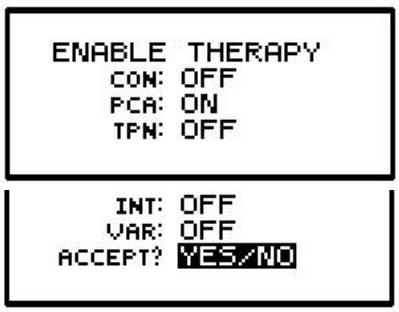
5 INVOICE NUMBER: This feature gives the programmer the option to use the numeric keypad to enter an invoice number (up to ten digits). If this option is not to be used, it can be left blank and skipped, using the down arrow to scroll to the next field. When the INVOICE NUMBER is satisfactory, press the "YES/ENTER" key at the "ACCEPT? YES/NO" field and the number will be stored in the pump history log.

6 CHANGE LOCK: This selection allows the clinician to change the lock setting of the pump. This feature is also available under the OPTIONS menu of each therapy. There are four choices in this field: OFF, 1, 2, or 3. When the lock is set to "OFF, " the clinician has access to all pump functions. Each level higher limits the options available to the patient or

caregiver (see each lock option screen in the individual therapy sections and/or the lock level table in the chapter entitled *Clinician Information and Access Code*.) If the desired lock does not appear on the display, press the "NO/CHANGE" key until the correct lock does appear. Then, press the "YES/ENTER" key to accept and move the cursor to the next field.

7 ENABLE THERAPY: This field is used to customize the **Curlin Infusion** pump to the specific therapy ordered for the patient. When you select this function by pressing the "YES/ENTER" key, the menu below appears, listing all five of the possible therapies. The pump's default setting is to enable PCA and disable all other therapies.

Enable Therapy Menu



You will not be allowed to change the default setting. If you attempt to enable a therapy, the following prompt will appear of this fact.

Patient History Log

The patient history log is a cumulative record of the approximately last 6,000 events that have occurred in the pump. Example: When prescription information is saved using the "YES/ENTER" key at the "DONE" or "ACCEPT" fields, a copy of the information also gets 'date and time stamped' and is recorded in the patient history log. Alarms, error codes, certain alerts, and certain key press events also get logged when they occur. The patient history log can be printed using the **Curlin Infusion** print package and by selecting the "Print Pt. Hx" option. Selecting the "Clear Hx Log" option will erase the patient history log. When the history log is full, it will overwrite on a "first-in, first-out" basis.

Note: The patient history log should be printed before clearing occurs if a hard copy record is required. The history log, when printed, lists the most recent event first. Clearing the history log does not affect the saved patient information.

When printed, the patient history log includes a heading at its beginning and a footer at its end. The heading includes the pump serial number, the patient history log title and revision number information of the software and hardware of the pump. The footer contains information about how much of the history log has been used. The information is printed with the most recent event first, and organized for easy transfer to a spreadsheet for sorting and reporting. A partial sample of a history log is given on the following page:

History Event Log

```
 SERIAL NUMBER              :201293
 MANUFACTURE DATE           :10AUG00
 MAINTENANCE DUE DATE       :16JAN04

 HISTORY EVENT LOG
22JAN03 15:38 ml/Vin/mmH/BV:9070.6/2.2/8/0.76
22JAN03 15:18 ml/Vin/mmH/BV:8937.3/2.2/10/0.76
22JAN03 14:58 ml/Vin/mmH/BV:8803.9/2.2/10/0.76
21JAN03 16:58 ml/Vin/mmH/BV:2.1/3.1/0/0.78
21JAN03 16:57 ml/Vex/mmH/BV:0.0/6.7/0/0.78
21JAN03 16:57 START PUMP   :0.0 ml
21JAN03 16:57 START THERAPY
21JAN03 16:57 AIR FILTER ? :YES
21JAN03 16:57 RUN/PAUSE KEY
21JAN03 16:57 CON RUN OPTION
              Options      :Lock OFF
              Lock         :OFF
              Hourly Tot   :OFF
              Clear Hrly   :OFF
              Dn Occlu     :HIGH
              Air Sens     :OFF
              Audio        :9
              PWR CK       :OFF
              Accept OPT   :YES/NO
21JAN03 16:57 CON Rx
              Bag Vol      :9900  ml
              Amt TBI      :9500  ml
              Rate         :400.0 ml/hr
              Time         :23:45 HH:MM
              KVO Rate     :0.1   ml/hr
              Done?        :YES
21JAN03 16:57 CON PRE Rx
              Units        :ml
              Delay        :OFF
              Titrate      :OFF
              Next?        :YES
21JAN03 16:57 REPEAT Rx
21JAN03 16:56 ON PUMP
21JAN03 16:33 OFF PUMP
Patient Hx Log Used (bytes): 1693 of 61440
```

Figure 4.1 History Event Log

8 PRINT PATIENT HISTORY: To print the patient history log, connect the Data Interface Cable to the communication port on left side of the pump before selecting this command. (FOLLOW DIRECTIONS THAT ACCOMPANY THE DATA INTERFACE CABLE.)

The pump will display:

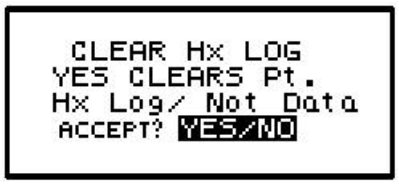
When you select "YES," the display will read "Printing Patient History Record." When the printing is completed, the pump display returns to the BIOMED SETUP Menu. To abort

the printing process, press the "NO/CHANGE" key on the pump's keypad, and the display will return to the BIOMED SETUP Menu. If the Patient History Log stops before all data is transferred, exit and restart the "Print Pt. Hx" command.

Note: To utilize the print feature of the **Curlin Infusion** pump, you must have a PC running the Hyperterminal program within MS Windows™, and use the **Curlin Infusion** Data Interface Cable which is purchased separately. Detailed user instructions are provided with the cable.

Clearing the History Log

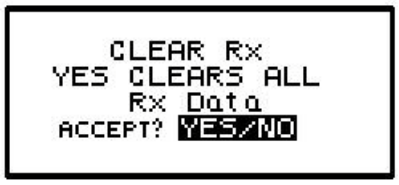
9 CLEAR Hx LOG: This feature is used to clear the patient history log from the pump. When you select "YES/ENTER," the pump displays:



When you select "YES," the history log is cleared and the display returns to the BIOMED SETUP Menu. Selecting "NO" aborts the feature and also returns to the menu.

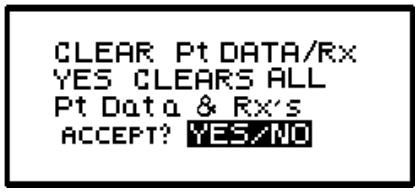
Clearing Patient Information

10 CLEAR Rx: When this command is selected, prescription information for all currently programmed therapies is cleared; however, the patient data is retained. When you select "YES/ENTER" the pump displays:



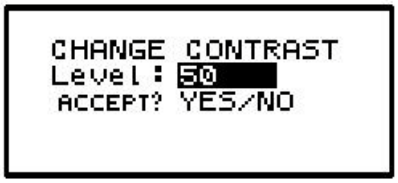
When you select "YES," all prescription programs are cleared, and the display returns to the BIOMED SETUP Menu. Selecting "NO" aborts the feature and also returns to the menu.

11 CLEAR Pt. DATA/RX: **Selecting this feature clears all data programmed and resets the pump to its default settings. The history log is NOT cleared.** When you select "YES/ENTER" at this field, the following screen displays:



When you select "YES," all patient data clears and the display returns to the BIOMED SETUP Menu. Selecting "NO" aborts the feature and also returns to the menu.

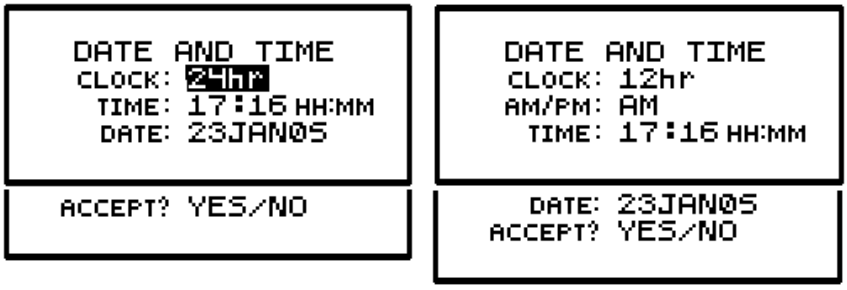
12 CHANGE CONTRAST: The contrast on the display screen is automatically optimized and is no longer changeable with this option.



Changing Date and Time

13 CHANGE DATE AND TIME: When this field is selected, the screen on the left appears:

Change Date and Time Screen



This screen is used to set the system Real Time Clock in the pump and will change the date and time displayed on the Welcome screen.

Note: The time must be reset to accommodate Daylight Savings Time changes and also at any time the pump is moved into a different time zone.

a. Time Format: You have the option of selecting twelve-hour or twenty-four-hour time format. Press the "NO" key until the correct selection appears on the field and then press the "YES/ENTER" key to accept the selection and move the cursor to the next field.

b. AM/PM: The second field allows you to select AM or PM when the twelve-hour time format is selected in the previous field. If the twenty-four-hour format was selected, this field will not appear. Press the "NO" key until the correct selection appears on the field, and then press the "YES/ENTER" key to choose the selection and move the cursor to the next field.

c. Time: Use the numeric keys to enter the correct time and then press the "YES/ENTER" key to accept and move the cursor to the next field.

d. Using the numeric keys, enter the correct mm/dd/yy, using two digits for each (e.g., if the month is January, enter 01; if it is a single digit date, use the number preceded by a 0 and then enter the last two digits of the year code, 00, 01, etc.). When the date is correct, press the "YES/ENTER" key to accept and move to the next field.

e. A final prompt will be given to "ACCEPT? YES/NO." If any data needs to be corrected, use the arrow keys to locate the correct field, press "NO" to change the data, enter the correction(s), and then press "YES/ENTER" again to accept the correction. Move the cursor back to the "ACCEPT? YES" field and press "YES/ENTER". This action enters the new time and date and moves to the next field in the BIOMED SETUP Menu. Pressing the "NO" key on this field aborts the change date and time selection and returns to the BIOMED SETUP Menu.

IMPORTANT INFORMATION

If a therapy is in progress, the date and time changing functions will not be accessible until the therapy is complete.

14 When BIOMED SETUP is complete, move the cursor to the DONE? Field and press the "YES/ENTER" key to return to the Program or Set-Up Menu Screen.

Program

Program/BioMed Setup Menu

To select "PROGRAM," use the arrow keys to move the cursor to the "PROGRAM" field and press the "YES/ENTER" key. If no therapy is already programmed, a "Select Therapy Screen" appears prompting you to select a therapy. If a therapy has been programmed, the following screen appears:

Resume/Repeat Rx/New Program Menu

There are three different ways that you can start a therapy:

- RESUME: Should be used if a therapy is interrupted. The pump picks up where it left off, and continues the therapy.
- REPEAT Rx: Should be used to hang a new bag and repeat the same therapy over again. The infusion history displayed is reset (volume infused, boluses requested, boluses received, etc). Shift Totals and Hourly Totals continue to accrue.
- NEW PROGRAM: Should be used to program a totally new Rx. All the therapy parameters in the pump are set to factory default and must be re-entered by the clinician.

The judicious choice of the option can help speed up programming. This screen gives you all three options:

Resuming a Therapy

1 RESUME—If a therapy is interrupted and the pump is turned off or paused before completion of the infusion, you may select "RESUME" by moving the cursor to this field and pressing the "YES/ENTER" key. A notification screen displays, and a prompt is given to run the pump. At this time, you can continue to infuse the prescription from the exact point at which it was interrupted.

Note: Patients should be taught proper time frames or limitations for leaving their IV bags or medication reservoirs "hanging" during an interrupted state and, if indicated, to change to a new bag according to the healthcare provider's policies and drug manufacturer's instructions.

Repeating a Pre-Programmed Therapy

2 REPEAT Rx—Use this feature if an infusion is complete and you wish to repeat the same prescription over again or if a therapy is programmed and the pump is turned off before the infusion is started. Select this field by moving the cursor to "REPEAT Rx," and press the "YES/ENTER" key.

If the pump has been programmed for more than one therapy and you wish to select a therapy at this time, move the cursor to "REPEAT Rx," and press the "YES/ENTER" key. This action will take you to the "Select Therapy Screen" and allow you to choose which of the programs you now wish to repeat.

Note: If you are choosing a therapy that is different than the last one selected, "Shift Totals" and "Hourly Totals" will be cleared.

If only one therapy is programmed and REPEAT Rx is selected, the cursor will automatically advance through the prescription menus. You can stop the auto-advance function by pressing any key (except ON/OFF) and then complete the confirmation of the prescription by pressing the "YES" key at the remaining fields.

Note: REPEAT Rx **does not allow any changes to the Pre-Prescription Menu.** If changes need to be made in Pre-Prescription selections (units, concentration, etc.) select "New Program" and reenter the entire prescription.

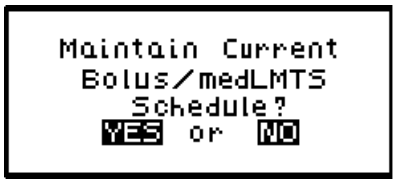
The lock settings affect the use of "REPEAT Rx". Lock 1 allows REPEAT with changes to the prescription screen only. Lock 2 allows REPEAT but does NOT allow the user to make changes in the prescription. Lock 3 does not allow REPEAT. (See lock setting table).

If a therapy has not completed and the REPEAT Rx function is selected, the following screen will display:

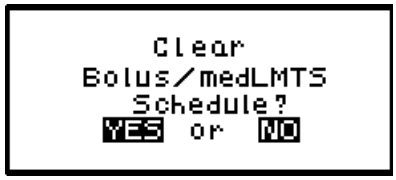
If the current infusion is to continue, press "NO," and you will be prompted to resume the

therapy.

If the current infusion is to be discontinued and a new IV bag started, press "YES/ENTER" to repeat. The following screen will then appear, asking whether or not you wish to maintain the current Bolus or MedLIMITS schedule.



Pressing "YES" will maintain the current schedules and continue with the programming of the pump. Answering "NO" will display the following screen, asking you to confirm the clearing of the schedules.



Pressing "YES" will clear the current schedules and continue with the programming of the pump. Answering "NO" will return you to the prior screen.

How to Go to "New Program"

NEW PROGRAM—If the cursor is moved to this field and the "'YES/ENTER" key is pressed, the display will automatically move to the first screen for a new programming.

Entering or Changing Data Fields

When prescription data fields appear, use the numeric keys to enter specific patient prescription information and, when correct, press the "YES/ENTER" key to accept. If a field on the menu is not "accepted" by the program, a context sensitive "Alert" prompt will appear and the cursor returns you back to the fields that require the change, to help you resolve the problem.

The following screen is the general alert prompt:

Programming a PainSmart pump with Protocol Library Safety System program

The information in this addendum describes how to program a pump with a drug protocol from the Protocol Library Safety System of a Curlin Infusion PainSmart™ (PCA only) Infusion Pump that has this program.

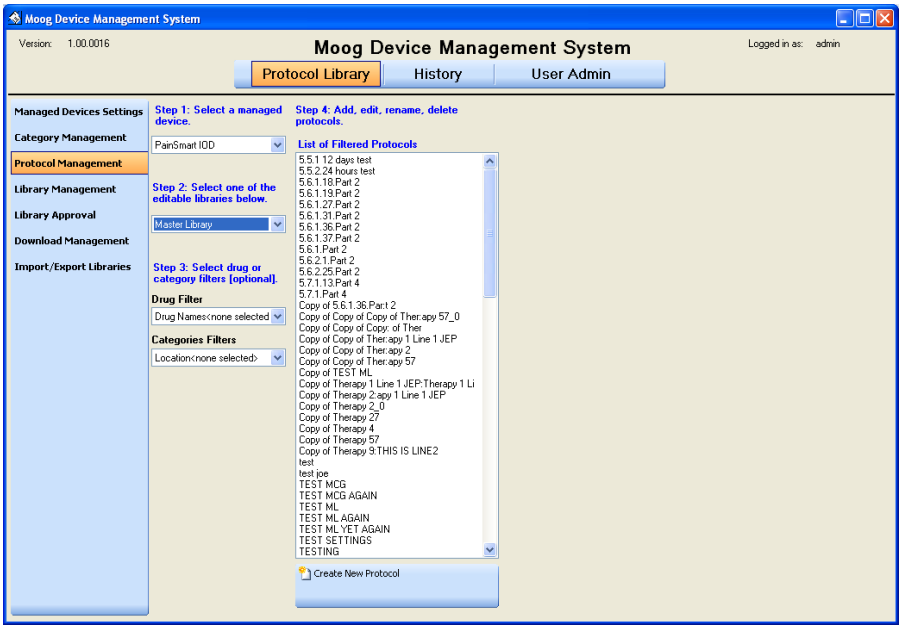
REQUIRED EQUIPMENT

The following equipment is required to service the pump:

- Data Interface Cable for PC (Curlin P/N: 340-2011)
- Personal Computer with Microsoft Windows: 2000, XP or Vista
- Protocol Library Safety System Software (Curlin P/N 360-7039)
- PainSmart™ IOD Ambulatory Infusion System with Protocol Library Safety System

Note: Electrostatic-Sensitive Devices, observe handling precautions.

Download Protocols from the Protocol Library Safety System Software into the Protocol Library-enabled Painsmart pump



How to navigate to the Download Management screen
Step 1: Click on "Protocol Library" from the top navigation bar.
Step 2: Click on "Download Management" from the left navigation bar.

How to connect the device to be downloaded
Step 1: Connect the data cable proper end to the device.
Step 2: Connect the RS232 end of the data cable to an available serial port on the desired PC.

How to download a library to the connected device
Step 1: Select the desired managed device from the "Managed Device" drop down.
Step 2: Select the proper comm. port from the "Managed Device Comm. Port" drop down menu.
Step 3: Select the desired library to download.
Step 4: Click on "Download Library" to start the download process.
Step 5: Immediately turn on the device that is connected to the PC.
Step 6: View the Download Status memo box to obtain the status of the download.

How to view or print the download history
Step 1: Click on "View Download Log" in order to view the download log.
Step 2: [Optional] Click on "Print Download Log" to print the log to the desired printer.
Step 3: Click on "Close Download Library" to revert to original screen.

Programming a Pump that has Downloaded Protocols in Place

YES key is used to verify or enter
NO key is used to change, erase or leave screen

Turn the Pump on, watch self test and power source bar.
Sets must be installed for pump to run.

Press the ⇑ and ⇓ keys to scroll through the menu, and press the **YES** key to accept/confirm a field entry and the **NO** key to reject/change a field entry.

1. Press ⇓ to scroll down to the LIBRARY function.
2. Highlight LIBRARY, and then press **YES**.
3. If Category/Care Area is not already listed on the first screen select **YES** to the right of "Page Down". This will bring up additional screen(s) with additional choices.

4. If Category/Care Area is already listed on first screen press ⇓ to scroll down to choice you wish to select.
5. When the Category/Care Area that you wish to choose is highlighted, press **YES** to select that option.
6. If the Drug Therapy is not already listed on the first screen select **YES** to the right of "Page Down". This will bring up additional screen(s) with additional choices.
7. If the Drug Therapy is already listed on first screen press ⇓ to scroll down to choice you wish to select.
8. When the Drug Therapy that you wish to choose is highlighted, press **YES** to select that option.
9. Press **YES** again to ACCEPT this Drug Therapy selection.
10. To use the drug Therapy as pre-programmed in the pump, select **YES** in each highlighted area to accept the information. To change any of the preselected drug therapy options when highlighted, press **NO** and enter correct information.

    Note: Some options may be "hard locked" so that changes cannot be made or there may be preset med limits that cannot be exceeded for safety.

Note: **YES** must be selected after each field when flashing to be accepted, if **YES** is not pressed after each field the pump will say: ALERT FIELD NOT ACCEPTED and will not allow you to continue programming.

Continue through each field until the RUN TO START screen appears after selecting done.

11. In the RUN TO START screen, you have three options:

    RUN TO START - to begin the infusion
    NO TO REVIEW - to check orders and make changes
    PRIME - to prime the tubing via the pump

12. In the RUN TO START screen choose **NO** to review, to validate therapy with another clinician. Press **YES** at each highlighted field to confirm choices, **NO** to change.
    (Note this should be done prior to priming tubing on the pump)

13. When you reach the RUN TO START screen for the second time you can now prime the tubing.
    To prime the tubing via the pump Press the PRIME button at the bottom of the pump, follow on screen instructions and hold the prime key down. Keep the button depressed until you are finished priming. Release the prime button to stop and press **YES TO EXIT** to return to the RUN screen.
14. You can now Begin your infusion by pressing **RUN TO START.**

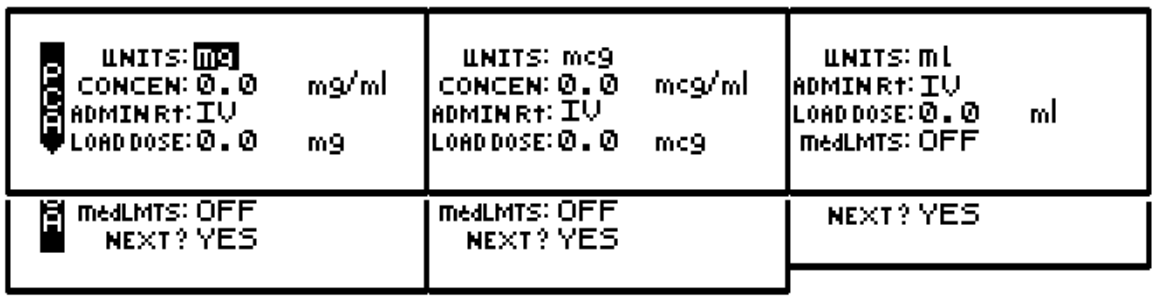
PCA IV Therapy

PCA or Patient Controlled Analgesia is designed for therapies requiring a continuous rate of infusion, patient controlled demand boluses, or both. This chapter covers Curlin Infusion pumps specifically programmed for PCA-Intravenous therapies.

When you begin programming a PCA-IV Therapy, the following display screen will appear, requiring you to enter information needed to program this therapy.

Prescription Menus

PCA-IV Pre-Prescription Menu Screen



These menu items allow you to customize the program to match each individual prescription. The pump's default settings are listed in the **left** screen. Each field is individually explained below.

Note: Some of the lines are abbreviated to fit on the display screen; the full descriptions appear below.

1 UNITS: Allows you to select from units of mg, mcg, or ml depending on how the prescription is written. In the PCA-IV therapy, the pump will allow rates as defined in figure 5.1; therefore, if mg or mcg is selected, the concentration requested in the next field will determine the upper and lower programming limits for mg and mcg. Use the "NO/CHANGE" key until the desired "units" appears on the display, and then press the "YES/ENTER" key to accept and move to the next field.

2 CONCEN (Concentration): This field is active whenever mg or mcg is selected in the "units" field and requires data entry of the concentration of mg or mcg per ml. When the correct data is entered, press "YES/ENTER" to accept, and move the cursor to the next field. If ml is chosen in the "units" field, this line will not display. The range for this field is 0.1 to 999.

3 ADMIN Rt (Administration Route): Allows a selection from three routes of administration: the default setting of IV (intravenous), EPI (epidural), and SQ (subcutaneous). If Epidural is selected, the Therapy Identifier bar in the Prescription Menu will display EPI instead of PCA. Each of the routes has defined settings as illustrated in the following table:

Administration Routes

Administration Route	Occlusion Setting mmHg (psi)	Maximum Basal Rate (ml/hr)	Rate During Bolus (ml/hr)	Maximum Bolus Dose (ml)	Maximum Load Dose (ml)
Intravenous	**900** or 400 (**18** or 8)	9.9	125	9.9	9.9
Epidural	**900** or 400 (**18** or 8)	25	90	25	25
Subcutaneous	**900** or 400 (**18** or 8)	5	60	5	5

Figure 5.1 PCA Administration Route Default Settings

4 Load Dose (Loading Dose): Allows the clinician to program in a pre-infusion loading dose if it is prescribed. If a loading dose is not ordered, select the default setting of "0" by pressing "YES/ENTER". If it is ordered, enter the dose in this field, and press the "YES/ENTER" key to accept and move to the next field. The maximum load dose is determined by the administration route (refer to Figure 5.1 above).

IMPORTANT INFORMATION

The amount of drug infused during the "loading dose" depends on the units chosen which may be in ml's, mg, or mcg. A loading dose of 5 mg will infuse 50 ml if the concentration is 0.1 mg/ml.

Note: The "units" will correspond to the choice previously selected.

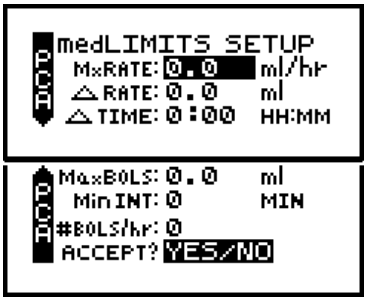
The loading dose will be delivered at the fixed bolus delivery rate when the "RUN" key is pressed to start the PCA infusion. The amount of the loading dose is included in the displayed "amount infused" and is logged into the history file. The bolus lockout time is set from the end of the loading dose. Because the loading dose is only used on initial start up of an infusion, it will revert to a value of "0" whenever the "REPEAT Rx" function is selected.

IMPORTANT INFORMATION

To terminate a loading dose in progress but save the rest of the prescription, pause the pump and select "REPEAT Rx." When the "RUN" key is pressed, the prescription will restart without the loading dose.

5 medLMTS™ Setup: Allows the clinician to setup soft (clinician or institutional) limits on any medication that is being infused with the pump. If there is no request or mandate to set soft limits, select "OFF," and press "YES/ENTER" to accept the factory defaults (see Figure 5.1). If a specific medication limit is requested or mandated, press "YES/ENTER" to turn ON the medLIMITS and the following screen will appear. Enter the appropriate soft limits for each field:

PCA medLIMITS™ Setup Screen



a. MxRATE (Maximum Rate) limits: Use the numeric keys to enter the upper limits of the infusion rate. When the correct maximum medLIMIT rate is entered, press "YES/ENTER" and the cursor will move to the next field. The minimum rate is 0, and the maximum rate must be equal to or less than maximum Basal Rate listed in Figure 5.1 Administration Routes table.

b. ΔRate (Delta Rate): Some prescriptions include incremental levels or *delta* limitations (e.g., "Do not raise the rate more than 1 ml in 12 hours"). If the prescription has these components, use the ΔRate field and the Δ Time field to enter those limits. Example: Enter the incremental medLIMIT amount level of 1 ml on this field, and press "YES/ENTER" to move to the next field.

c. ΔTime (Delta Time): Enter the incremental or delta time limit. (Example: Enter 12:00 hours.) If there are no incremental or delta level components to the prescription, leave these fields (ΔRate and ΔTime) set at their default settings of "0" and press "YES/ENTER" to move to the next field.

d. Max BOLS (Maximum Bolus Dose): Use this field to set the upper limit of the Patient Bolus or Patient Demand dose. If there is no bolus dose, enter "0." If a value is given for the Bolus Dose, use the numeric keys to enter it on this field, press the "YES/ENTER" key to accept, and move to the next field. The value entered for this field must be equal to or less than maximum Bolus Dose listed in Figure 5.1 Administration Routes table.

e. MinINT (Minimum Bolus Interval): Use this field to set the minimum amount of time allowed between patient bolus doses. When the correct time is entered, press the "YES/ENTER" key to accept, and move to the next field.

f. #BOLS/hr (Number of Boluses allowed per hour): Use this field to enter the maximum patient bolus/hr. When correct, press the "YES/ENTER" key to accept, and move to the next field.

g. ACCEPT? If there are any changes to be made, use the up or down arrows to move to that field, use the numeric keys to enter the corrected data, and press "YES/ENTER" to accept the new values. Move the cursor back to the "ACCEPT?" field and press "YES/ENTER". The medLIMITS are now entered and the display returns to the Pre-Prescription screen. If the entire medLIMITS Setup Screen is to be aborted, press "NO" to return to the Pre-Prescription screen.

6 NEXT? If any field is not satisfactory, use the arrow keys to locate the field to be changed, enter the changes, press the "YES/ENTER" key to accept the new values and then go back

to the "NEXT?" field. When all fields are satisfied with correct settings, press the "YES/ENTER" key to exit this screen, and move on to the next PCA Programming Menu.

At the following screen, the patient's prescription information is entered to complete the programming of the pump for a PCA Therapy. Remember that there may be additional limits to rates and doses based on limits for the route of administration selected. (See Figure 5.1).

Note that the screen Identifier gives the abbreviation "PCA" for Patient Controlled Analgesic Therapy (or EPI if Epidural route is selected) and has a down arrow configuration, which indicates that there are more than four fields of text or information to be considered.

PCA Prescription Menu

Note: The PCA therapy can be programmed for Basal Rate (continuous infusion) only, for Patient Bolus (patient demand dose) only, or for both. At least one of these settings must be entered, though, before this menu can be accepted at the "DONE?" prompt.

1 BAG Vol (IV Bag Volume): Using the numeric keys, enter the amount of fluid in the IV Bag or Medication Reservoir (include the overfill volume).

Note: The volume used for priming via the pump is deducted from the Bag Volume. Therefore, Bag Volume should be equal to or greater than the Amount To Be Infused to start the infusion.

When the correct amount has been entered, press the "YES/ENTER" key to accept this field and move on to the next. The range for this field is 1 to 9999.

2 BASL RATE (Basal Rate): The **Curlin Infusion** pump in PCA Intravenous Therapy can be set for a continuous basal infusion (see Figure 5.1). Use the numeric keys to enter the prescribed rate (the "units" chosen in the Pre-Prescription Menu will appear here).

Note: If the rate falls out of the range of the pump (depending on the administration route selected), a beep will sound, and one of the following alert messages will appear:

This field will not be accepted with the "YES/ENTER" key until the rate is adjusted to fall within pump's range. When the rate is correct, use the "YES/ENTER" key to accept, and move to the next field.

3 Pt. BOLUS (Patient Bolus Dose): Use the numeric keys to enter the prescribed bolus (patient demand dose) and when correct, press the "YES/ENTER" key to accept and move on to the next field. (see Figure 5.1).

Note: If a bolus dose entered falls out of the range of the pump, the above "Greater than Max Field Limit" or "Less than Min Field Limit" alert will also appear and display the limit.

4 BOLS INT (Patient Bolus Interval): This field allows the clinician to enter the minimum amount of time (from 0 to 60 min) prescribed between patient-activated bolus dosing. Enter the time using the numeric keys and, when correct, press the "YES/ENTER" key to accept and move the cursor to the next field.

5 # BOLS/hr (Number of Patient Bolus Doses per hour): Using the numeric keys, enter the number (0 to 15) of boluses allowed per hour and, when correct, press the "YES/ENTER" key to accept and move to the last field.

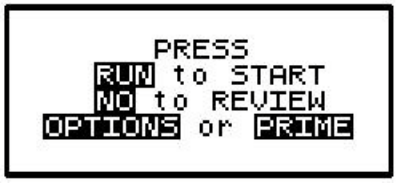
6 DONE?: If the programming is not done, use the up or down arrow keys to locate the field to be changed, use the numeric keys to enter the corrected information and press "YES/ENTER" to accept the new data. If all the fields are acceptable, at the "DONE?" prompt, press the "YES/ENTER" key to move on to the next screen.

IMPORTANT INFORMATION

Before the infusion is started, it is important to consider a few remaining issues, such as checking the administration set to be sure it was primed or purged of all air, or perhaps the pump may need to be put into a security lock setting for additional safety.

The following screen offers you four choices:

Run Options Screen



Note that you do not need to answer yes or no questions or enter data as previously requested. This screen requires only that you press the corresponding key on the keypad to begin the requested function. The following section explains each selection beginning with Prime:

Prime

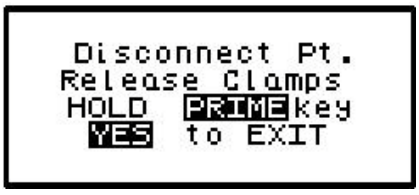
The IV bag or syringe and administration set should be inspected and be free of all air and air bubbles before they are connected to the access site. Priming may have been done by gravity when the administration set was first connected to the bag (see Chapter 2). However, if it was not done then or if there are any remaining air bubbles to be removed, the pump's "PRIME" function will assist in this process. To prime the set by using the pump, follow the directions below.

Note: The lock setting may restrict the ability to prime using the pump.

1 Press and release the "PRIME" key at the "Run Options" screen.

2 The following screen appears:

Prime Direction Screen



IMPORTANT INFORMATION

3 Disconnect the administration set from the patient's access site using aseptic protocols.

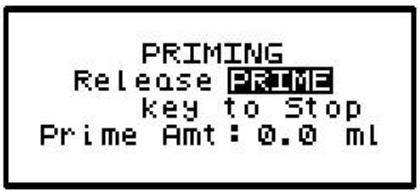
Priming with the set connected to the patient could result in overdose and may cause injury or even death to the patient.

4 Release any clamps on the administration set.

5 Press and hold the "PRIME" key. Fluid will be pumped at a rate of 300 ml/hr through the administration set as long as the PRIME key is pressed or until 6 ml is pumped. The pump will then beep once and require repressing and holding the "PRIME" key again until 6 more milliliters is delivered. Continue this process until all air has been purged from the IV bag and the administration set.

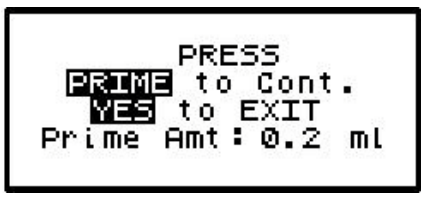
Note: If the bag volume is depleted during priming, a subsequent priming event cannot be initiated.

While the "PRIME" key is depressed, the following screen appears:



The amount used for the prime session is accumulated and displayed on the bottom field of this screen.

6 When priming is complete, release the "PRIME" key. The following screen will display:



7 To continue priming, press the PRIME key again. If priming is complete, press "YES/ENTER" to exit and return to the Run Option screen. The total amount of solution used in this priming session displays on the bottom field.

Note: The amount of fluid used in the priming process will be deducted from the Bag Volume Total but will NOT be counted in the amount infused because this amount is not to be delivered to the patient. The bag volume should still be equal to or greater than the Amount TBI before you can proceed with the infusion. If it is not, an alert will appear stating, "AMT TBI GREATER THAN BAG VOL". You have the option to continue by pressing the "YES/ENTER" key or return to the "Run Options" screen by pressing the "NO" key.

Options

Once the prescription is programmed, there are more customizing features available for each therapy. Press the "OPTIONS" key from the Run Options screen to access these options. The OPTIONS menu can also be accessed while the pump is running.

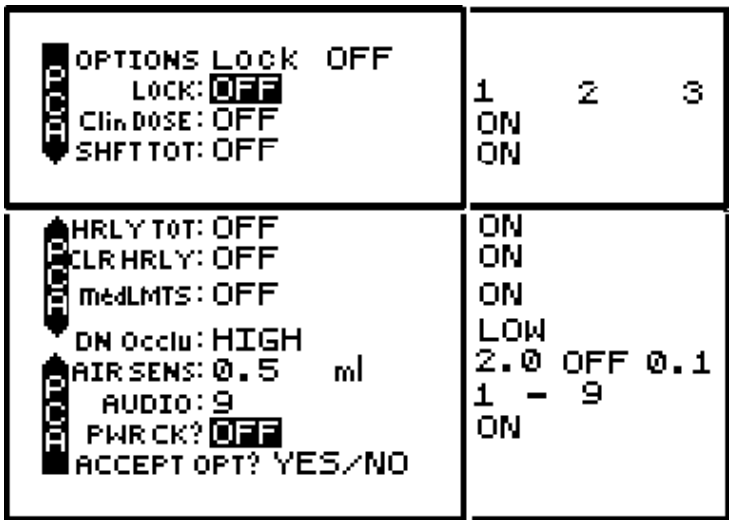
Note. After the infusion has started, any changes made through the OPTIONS menu involving:

- "Clin DOSE",
- "SHFT TOT"
- "HRLY TOT", and
- "CLR HRLY"

will return you directly to the "ACCEPT OPT?" field.

The following screen displays the PCA Therapy "Options" available. When you open this menu, use the up and down arrows to move the highlight bar to the desired option. Lock settings may require an access code before the option can be changed. Any new setting, except "CLEAR HOURLY" screen acceptance and "SHIFT TOTAL" clear screen acceptance becomes effective when the "Options" screen is exited by pressing "YES/ENTER" at the "ACCEPT OPT?" field.

PCA Options Menu



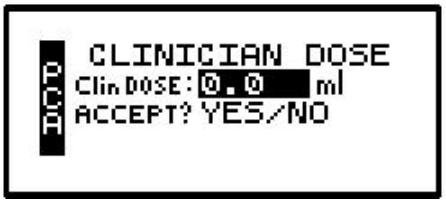
Note: The pump's default settings are indicated above in the **left** screen.

1 LOCK: When the lock is set to "OFF," the clinician has access to all of the pump functions. After the pump is completely programmed, change the lock to meet the security needs of this therapy. There are four choices: OFF, 1, 2, or 3. Each level higher limits the available options. To change the lock setting, press the "NO/CHANGE" key until the correct lock setting appears. Then, press "YES/ENTER" key to accept the new lock selection. Once the pump has been set into a locked condition, any subsequent request to change the lock will

require a clinician access code. (See "Clinical Information and Access Code" chapter for access code information, lock setting table and available features for each lock setting).

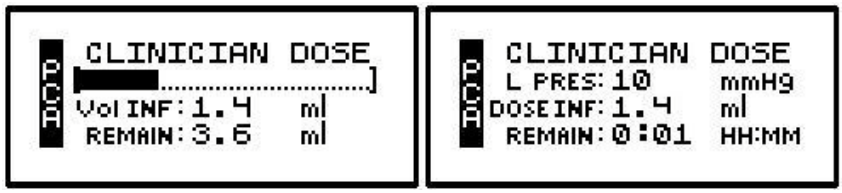
2 ClinDOSE? (Clinician Dose): Allows an additional prescribed dose to be administered by a clinician at any time except during a Load Dose or Bolus Dose. The Maximum Clinician Dose is determined by the administration route (see Figure 5.1).

Clinician Dose Menu



Note: If a lock setting of 1, 2 or 3 is selected, an access code will be requested. If you wish to give a clinician dose at this time, change OFF to ON and press YES key. Enter the amount of clinician dose to be delivered. Press YES to accept the clinician dose menu. The Clinician Dose will begin when the options screen is accepted and the pump is running. The following Run Screen will appear until it is completed:

Clinician Dose Run Screen



The graphic bar now depicts the clinician dose in progress and when this dose is complete, reverts back to reflecting the container in progress. **This graphic bar will toggle with Line Pressure ("L Press") which displays the current patient side (downstream) pressure in the administration set in millimeter of mercury (mmHg) units.**

The Volume Infused will toggle with Dose Infused if mg or mcg are the selected unit. Remaining dose time will toggle with remaining dose volume to infuse.

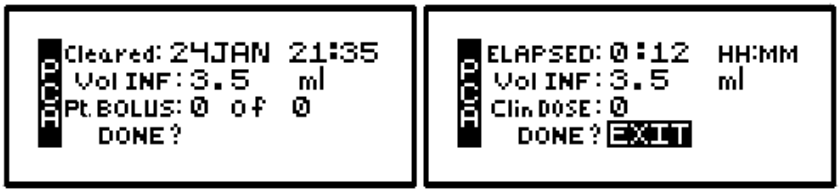
IMPORTANT INFORMATION

When the clinician dose is complete, the pump will beep, return to the previous running rate, and resume the PCA therapy. <u>The lockout time for bolus dosing will restart at the end of the clinician dose to prevent a bolus from being delivered too quickly after a clinician dose</u>.

Once the clinician dose is given, the clinician-dose field returns to its default of "0," and any subsequent clinician doses will need to be reentered.

3 SHFT TOT (Shift Totals): This feature allows the clinician to track the infusion for a selected period of time, such as an 8 or 12-hour shift. To access Shift Totals, press the "NO/CHANGE" key to change OFF to ON and then press "YES/ENTER." The following screens will appear alternately:

Shift Totals Screens



a. Cleared/ELAPSED: This field displays the date and time the shift total was last cleared. It toggles with the amount of time elapsed since that clearing.

b. Vol INF/Amt INF (Volume/Amount Infused): This field displays the volume (ml) and amount (mg or mcg) infused since the shift total was cleared.

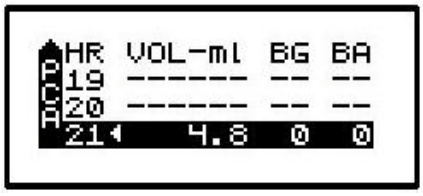
c. BOLUS/Clin DOSE (Clinician Dose): This field displays the number of clinician doses given since the total was cleared. It toggles with the number of patient activated boluses received of those requested.

d. To exit the shift total feature without clearing the totals, press the "YES" key at the highlighted EXIT field.

e. To clear the Shift Totals, press the "NO" key at the EXIT field. The field will change to read "CLEAR", press the "YES" key now to clear. A lock level 3 will require an access code entry before allowing the user to clear the Shift Totals.

4 HRLY TOT (Hourly Totals): To view hourly infusion totals press the "NO/CHANGE" key to change OFF to ON and then press "YES/ENTER." Either of the following two screens will appear:

Hourly Totals Screen



or

The screen displays hourly infusion delivery for each hour (00-23) of the last 24 hours of therapy in the unit selected.

a. The highlighted field has a marker ◄ indicating the current hour in progress.

b. Volume or Amount will show in the first column, BG (Bolus Given) in the second column and BA (Bolus Attempted) in the third column.

c. Use the up arrow key to scroll back through the last 24 hours.

d. If the infusion was not running, a series of ------ will appear for that hour.

e. To exit this screen, press the NO/CHANGE key.

5 CLR HRLY (Clear Hourly Totals): To clear Hourly Totals press the "NO/CHANGE" key to change OFF to ON and then press "YES/ENTER." The following screen will appear:

Clear Hourly Totals Screen

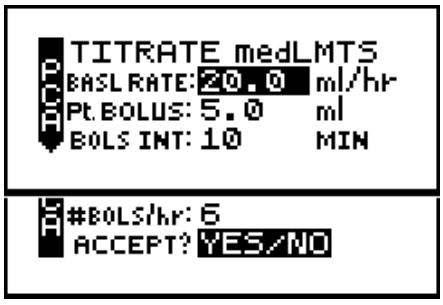
a. To clear the Hourly Totals, press the "YES/ENTER" key. All Hourly Totals will be cleared and the cursor will return to the Options menu.

b. To maintain the Hourly Totals, press the "NO/CHANGE" key. Hourly Totals will be maintained and the cursor will return to the Options menu.

c. A lock level of 2 or 3 will require an access code entry before allowing "Hourly Totals" reset.

6 medLMTS™ (Medication Limits): If this feature was **not** selected in the Pre-Prescription Menu, it will not appear on the Options menu. If medLIMITS™ parameters were programmed previously, this field will appear; and if you want to titrate now, select "ON" and press the "YES/ENTER" key to go to the PCA Titrate medLIMITS™ menu:

PCA Titrate medLIMITS™ Screen



This screen allows you to change the basal rate and/or the bolus parameters within the predefined prescription limits. Whenever any value is changed using the medLIMITS™ option, the new value will appear on the prescription menu when it is reviewed. If a delta time/rate limitation was previously programmed, the rate can be increased within those parameters up to 12 times during the delta time period. If a 13th attempt is made an alert will display: "MAX CHANGES FOR PERIOD REACHED."

a. BASL RATE: Use the numeric keys to enter the new desired rate. When the correct rate has been entered, press the "YES/ENTER" key to accept the rate, and move to the next field. (see Figure 5.1).

b. Pt. BOLUS: Use the numeric keys to enter the new patient bolus dose and, when correct, press "YES/ENTER" to accept and move to the next field. (see Figure 5.1).

c. BOLS INT (Patient Bolus Interval): Use the numeric keys to enter the new patient bolus time interval and, when correct, press "YES/ENTER" to accept and move to next field. The range for this field is 0 to 60.

d. #BOLS /hr (number of patient boluses per hour): Use the numeric keys to enter the new number of patient boluses per hour and, when correct, press "YES/ENTER" to move to the final step in the medLIMITS menu. The range for this field is 0 to 15.

e. ACCEPT? YES/NO: If necessary, use the arrow keys to move to any field and change the data entered. To abort this screen entirely, press "NO" at this prompt. Any data entered on this display will be aborted and the display will return to the options menu . If the parameters entered are correct, press the "YES/ENTER" key, and the new rate or rates will be accepted. The display returns to the Options menu when this screen is exited.

f. New values will be in effect when the Options menu is accepted.

7 DN Occlu (Down Occlusion): This feature allows the pump to sense an occlusion pressure in the line between the pump and the access site, e.g., a kink in the tubing between the pump and the patient, a blockage in the access site, or a clamp left on the distal tubing. There are two options for this field: the "Low" setting triggers the alarm when it detects in-line pressures at 400 mmHg ±150 (8 ±3 psi "pounds per square inch"), or the "High" setting (default setting) alarms when it detects in-line pressures at 900 mmHg ±150 (18 ±3 psi). Press "NO" to change the selection. When the correct pressure setting appears, press the "YES/ENTER" key to accept it and move the cursor to the next field.

8 AIR SENS (Air In Line Sensor): This feature allows you to select the amount of air the pump will detect in the administration set tubing before it goes into an alarm state. The choices are 0.1 ml, 0.5 ml (the default setting), 2.0 ml, and OFF. Press the "NO/CHANGE" key to change the options until the desired setting appears on the display, and then press "YES/ENTER" to select and move the cursor to the next line.

IMPORTANT INFORMATION

Whenever the Air-In-Line Sensor is set to OFF, Curlin Infusion recommends the use of an air-eliminating filter.

When the Air-In-Line Sensor is set to "OFF" and the "RUN" key is pressed at the Run Option screen, the alert message below will appear on the screen. It will also appear if the Air-In-Line is changed from the OPTIONS menu when an infusion is in progress and the AIR SENS field is accepted.

Air-In-Line Sensor Off Message Screen

This message alerts you to the Air-In-Line Sensor being set to "OFF." The message will require answering "yes" or "no" to whether the administration set being used has an air-eliminating filter. If you press the "YES" key to indicate that there is an air-eliminating filter in use, the Air-In-Line sensor will be completely disabled and the pump will start the infusion.

If you press the "NO" key to indicate a filter is <u>not</u> in use, the Air-In-Line sensor will be set to detect the presence of any contiguous 2-milliliter air in the line. The alert message below will then briefly appear on the screen. If the alert occurred at the Run Options screen, the pump will start the infusion.

Pump Will Sense 2ml Alert Screen

Also, when the Air-In-Line sensor is completely disabled and the infusion goes to an "Infusion Complete" or "Empty Bag" status, the Air Sensor setting will automatically change to 2.0 ml.

9 AUDIO (Audio Volume): This feature allows you to adjust the audio volume of the pump's alarm setting. You can enter a number from 9 (the loudest setting) to 1 (the quietest setting). When you select a number, the pump will beep at that audio level. Press the "NO" key and enter a new setting (1 – 9). Repeat until the desired setting is reached, and then press "YES/ENTER" to select and move the cursor to the next field.

10 PWR CK? (Power Check): This feature allows you to periodically recheck the battery charge. It will display the active power source of the pump, whether internal batteries or external power. If the active source is the batteries, a graph will indicate approximately how much charge remains in the batteries. To access this information, select "ON" and press "YES/ENTER". The Power Source Screen will display for five seconds and return to the Options Menu.

11 ACCEPT OPT? If the changes made in the Options Menu are acceptable, press the "YES/ENTER" key. If the changes are not acceptable, cursor up to any field for corrections or abort all changes made in this menu by pressing the "NO" key, and return to the previous screen.

There are two remaining selections to consider from the "Run Options Screen", **NO** to **REVIEW** and **RUN** to **START**.

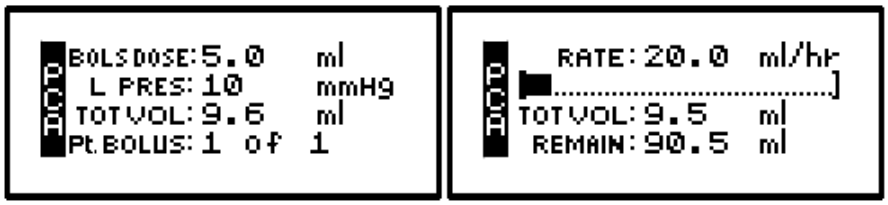
Reviewing or Changing the Program

To review or make any changes in the prescription programming, press the "NO/CHANGE" key at the Run Options screen, and the prescription screens will reappear (Changes may be inhibited by lock setting.) Changes to the prescription can be made before the start of the therapy; however, once a therapy has started, no changes can be made to the Pre-Prescription Screen. It will appear only for review purposes. Attempts to change the Pre-Prescription Menu during a new program review will cause an alert screen to be displayed, notifying the programmer that changes will require reprogramming of the prescription.

Starting the Infusion

RUN: When the programming is complete and the pump has been primed properly and connected to the access site, press the "RUN" key at the Run Option screen to start the therapy. The pump will begin infusing, and except when a LOAD DOSE or CLIN DOSE is programmed, the following screen will display data on the status of the therapy:

PCA Run Screen



IMPORTANT INFORMATION

The amount of drug infused during the "loading dose" depends on the units chosen which may be in ml's, mg, or mcg. A loading dose of 5 mg will infuse 50 ml if the concentration is 0.1 mg/ml.

This screen will continue to display as long as the infusion is running. Notice the information on each field:

1 The first field displays the "RATE" of the infusion toggled with the programmed Patient Bolus Dose Amount. This field also displays "BOLUS xx (mg/ml/mcg)" whenever a bolus dose is being delivered. The programmed bolus amount displays and the BOLUS text flashes when the pump is delivering a patient bolus dose.

2 The second field is a graphic bar representing the percentage of the completed infusion. **The graphic bar will toggle with Line Pressure ("L Press") which displays the current patient side (downstream) pressure in the administration set in millimeter of mercury (mmHg) units.**

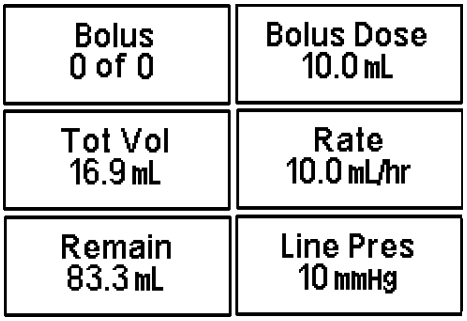
3 The third line displays the total volume "TOT Vol" infused and, if mg or mcg is the selected unit, this field will toggle to display the total mg or mcg infused "TOT INF" as well.

4 The fourth line displays the number of patient boluses delivered and number of boluses attempted per bag volume. If the patient attempts a bolus before the allowed amount of time set between boluses, an alert message will display "Bolus Requested," and only the second number will increase. Also, toggling with number of boluses delivered/requested is the remaining volume left in the bag.

5 In addition to the display screen, information on the status of the infusion is provided by the three LED lights. The green LED light blinks whenever the pump is infusing. If the Basal rate is set at "0", the yellow LED light will blink along with the green light until the bolus is actually being delivered. While the bolus is infusing, only the green light will blink.

The information on the run screen will alternate between a four-line display and a large font display of each individual line of information on the screen.

PCA Information Screens



Interrupting an Infusion

The infusion can be paused at any time by pressing the "PAUSE" key. The yellow "Standby" LED light blinks, and the following screen displays:

Pause Menu Screen

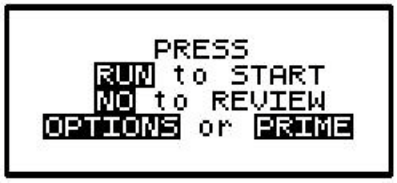
The screen offers the user three options:

Resuming the Current Container

1 RESUME: To resume the therapy in progress from where it left off, press the "YES/ENTER" key at this field. The pump will return to the Run Options Screen, and you can resume the therapy.

Note: If you wish to hang a new container, move the cursor to REPEAT Rx (see following section).

Run Options Screen



a. Press the "RUN" key to resume the therapy.

b. Press the "NO" key to go back to the programming screen to make changes in the therapy program (this feature may be blocked by a security lock, thus not allowing changes to the program unless an access code is given). Also, remember that no changes are allowed in the first prescription menu if an infusion is in progress.

c. Press the "OPTIONS" or "PRIME" keys if either of these functions is desired at this time.

Hanging a New Container

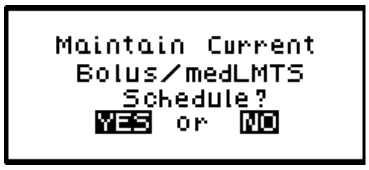
2 REPEAT Rx: To repeat the present therapy and add another IV container with the same prescription, press the "YES/ENTER" key at this field. If more than one therapy is programmed, a "Select Therapy Screen" will list all preprogrammed therapies. Select which therapy to repeat by moving the cursor to the therapy of choice and pressing the "YES/ENTER" key.

The pump will then display the program screen with the last programmed values of the therapy and the cursor will automatically advance through the prescription menus. You can stop the auto-advance function by pressing any key (except ON/OFF) and then complete the confirmation of the prescription manually by pressing the "YES" key at the remaining fields.

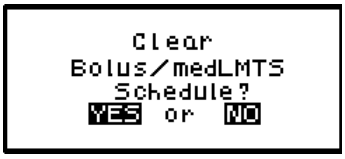
When the program is accepted, the display returns to the above "Run Options Screen." If the current therapy is still in progress when the "REPEAT Rx" function is selected, the following screen will appear:

a. This screen is a reminder that the infusion is not complete. If the incomplete therapy is to be continued, press "NO/CHANGE" and the Pause Menu screen will be displayed, then press "YES/ENTER" and the Run Options screen will be displayed. To resume the unfinished therapy, press "RUN".

b. If the therapy is to be terminated and a new IV container added, press "YES/ENTER" to repeat the infusion. The following screen will then appear, asking whether or not you wish to maintain the current Bolus or MedLIMITS schedule.



Press "YES/ENTER" to maintain the current schedules and continue with the programming of the pump. Press "NO/CHANGE" to start a new schedule. The following screen will then appear asking you to confirm the clearing of the schedules.



Press "YES/ENTER" to clear the current schedules and continue with the programming of the pump. Pressing "NO/CHANGE" will return you to the prior screen.

Entering a New Program

3 NEW PROGRAM: To program an entirely new prescription or different therapy, press the "YES/ENTER" key at this field (the ability to change these parameters may be controlled by the lock setting). The display will return to the "Pre-Prescription Screen." If the current container has not completely infused, the following screen will appear:

a. This screen is a reminder that the infusion is not complete. If you want to return to the current container and finish infusing it, press the "NO/CHANGE" key to return to the Pause Menu screen.

b. To abort the current container and enter a new program, press the "YES/ENTER" key. If a prescription has previously been entered for the therapy mode you wish to program, the following screen will appear:

c. If you want to save the current prescription, press the "NO/CHANGE" key to return to the Pause Menu screen. If you wish to erase the current prescription and enter a new program, press the "YES/ENTER" key to proceed. All previous prescription values will be erased.

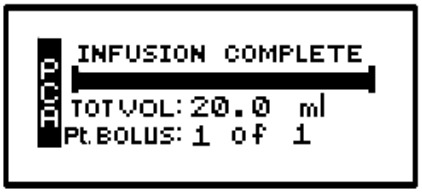
Note: Selecting "NEW PROGRAM" will also reset the OPTIONS selected to their default settings except for the LOCK and AUDIO settings which must be changed manually.

Infusion Complete

When the infusion is complete, the audio alarm will beep, and the following screen will display:

Note: To temporarily silence the alarm, press the "Silence" key.

PCA Infusion Complete



This screen displays:

1 "INFUSION COMPLETE."

2 A completely filled in bar graph indicating infusion complete.

3 TOT Vol (Total Volume): If units are mg or mcg, this field will also toggle to give the Total Infused.

4 Infusion Complete is considered an alarm status, and the audio alarm will beep periodically every three seconds to alert the user to this condition. Additionally, the red LED light blinks to indicate that the "Infusion Complete" alarm exists.

Stopping the Infusion

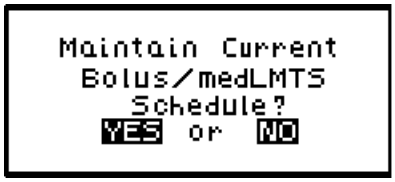
At "Infusion Complete", press the "PAUSE" key and then choose to either:

1 Turn off the pump by pressing the "OFF" key, or

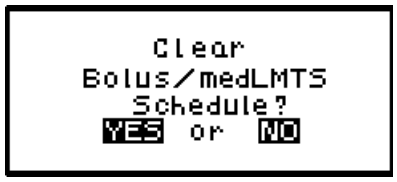
2 Select one of the options from the "Pause" screen:

Pause Menu Screen

a. REPEAT Rx: To repeat the present therapy and add another IV container with the same prescription, press the "YES/ENTER" key at this field. The following screen will then appear, asking whether or not you wish to maintain the current Bolus or MedLIMITS schedule.



Press "YES/ENTER" to maintain the current schedules and continue with the programming of the pump. Press "NO/CHANGE" to start a new schedule. The following screen will then appear asking you to confirm the clearing of the schedules.



Press "YES/ENTER" to clear the current schedules and continue with the programming of the pump. Pressing "NO/CHANGE" will return you to the prior screen.

The pump will then display the program screens with the last programmed values of the therapy and the cursor will automatically advance through the prescription menus. You can stop the auto-advance function by pressing any key (except ON/OFF) and then complete the confirmation of the prescription manually by pressing the "YES" key at the remaining fields.

b. NEW PROGRAM: To program an entirely new prescription press the "YES/ENTER" key when this field is highlighted and the display will return to the "Select Therapy Screen." (The lock setting can control access to this function.)

Note: The therapy is complete; therefore, RESUME is not an option at this time and is not displayed.

Chapter 6

PCA Epidural Therapy

PCA or Patient Controlled Analgesia is designed for therapies requiring a continuous rate of infusion, patient controlled demand boluses, or both. This chapter covers ***Curlin Infusion*** *pumps specifically programmed for PCA-Epidural therapies.*

When you begin programming a PCA-EPI Therapy, the following display screen will appear, requiring you to enter information needed to program this therapy.

Prescription Menus

EPI Pre-Prescription Menu Screen

UNITS: mg CONCEN: 0.0 mg/ml ADMIN Rt: EPI LOAD DOSE: 0.0 mg	UNITS: ml ADMIN Rt: EPI LOAD DOSE: 0.0 ml medLMTS: OFF	UNITS: mcg CONCEN: 0.0 mcg/ml ADMIN Rt: EPI LOAD DOSE: 0.0 mcg
medLMTS: OFF NEXT? YES	NEXT? YES	medLMTS: OFF NEXT? YES

These menu items allow you to customize the program to match each individual prescription. The pump's default settings are listed in the **left** screen. Each field is individually explained below.

Note: Some of the lines are abbreviated to fit on the display screen; the full descriptions appear below.

1 UNITS: Allows you to select from units of mg, mcg, or ml depending on how the prescription is written. In the PCA-EPI therapy, the pump will allow rates as defined in figure 5.1; therefore, if mg or mcg is selected, the concentration requested in the next field will determine the upper and lower programming limits for mg and mcg. Use the "NO/CHANGE" key until the desired "units" appears on the display, and then press the "YES/ENTER" key to accept and move to the next field.

2 CONCEN (Concentration): This field is active whenever mg or mcg is selected in the "units" field and requires data entry of the concentration of mg or mcg per ml. When the correct data is entered, press "YES/ENTER" to accept, and move the cursor to the next field. If ml is chosen in the "units" field, this line will not display. The range for this field is 0.1 to 999.

3 ADMIN Rt (Administration Route): Allows a selection from three routes of administration: the default setting of EPI (epidural), IV (intravenous) and SQ (subcutaneous). If a route other than Epidural is selected, the Therapy Identifier bar in the Prescription Menu will display PCA instead of EPI. Each of the routes has defined settings as illustrated in the following table:

Administration Routes

Administration Route	Occlusion Setting mmHg (psi)	Maximum Basal Rate (ml/hr)	Rate During Bolus (ml/hr)	Maximum Bolus Dose (ml)	Maximum Load Dose (ml)
Epidural	**900** or 400 (**18** or 8)	25	90	25	25
Intravenous	**900** or 400 (**18** or 8)	9.9	125	9.9	9.9
Subcutaneous	**900** or 400 (**18** or 8)	5	60	5	5

Figure 6.1 PCA Administration Route Default Settings

4 Load Dose (Loading Dose): Allows the clinician to program in a pre-infusion loading dose if it is prescribed. If a loading dose is not ordered, select the default setting of "0" by pressing "YES/ENTER." If it is ordered, enter the dose in this field, and press the "YES/ENTER" key to accept and move to the next field. The maximum load dose is determined by the administration route (refer to Figure 6.1 above).

IMPORTANT INFORMATION

The amount of drug infused during the "loading dose" depends on the units chosen which may be in ml's, mg, or mcg. A loading dose of 5 mg will infuse 50 ml if the concentration is 0.1 mg/ml.

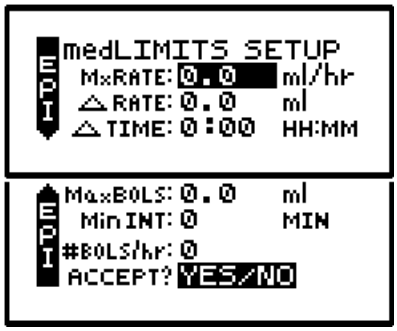
Note: The "units" will correspond to the choice previously selected.

The loading dose will be delivered at the fixed bolus delivery rate when the "RUN" key is pressed to start the PCA infusion. The amount of the loading dose is included in the displayed "amount infused" and is logged into the history file. The bolus lockout time is set from the end of the loading dose. Because the loading dose is only used on initial start up of an infusion, it will revert to a value of "0" whenever the "REPEAT Rx" function is selected.

IMPORTANT INFORMATION

To terminate a loading dose in progress but save the rest of the prescription, pause the pump and select "REPEAT Rx." When the "RUN" key is pressed, the prescription will restart without the loading dose.

5 medLIMITS™ Setup: Allows the clinician to setup soft (clinician or institutional) limits on any medication that is being infused with the pump. If there is no request or mandate to set soft limits, select "OFF," and press "YES/ENTER" to accept the factory defaults (see Figure 6.1). If a specific medication limit is requested or mandated, press "YES/ENTER" to turn ON the medLIMITS and the following screen will appear. Enter the appropriate soft limits for each field:



a. MxRATE (Maximum Rate) limits: Use the numeric keys to enter the upper limits of the infusion rate. When the correct maximum medLIMIT rate is entered, press "YES/ENTER" and the cursor will move to the next field. The minimum rate is 0, and the maximum rate must be equal to or less than maximum Basal Rate listed in Figure 6.1 Administration Route Default Settings table.

b. ΔRate (Delta Rate): Some prescriptions include incremental levels or *delta* limitations (e.g., "Do not raise the rate more than 1 ml in 12 hours"). If the prescription has these components, use the ΔRate field and the Δ Time field to enter those limits. Example: Enter the incremental medLIMITS amount level of 1 ml on this field, and press "YES/ENTER" to move to the next field.

c. ΔTime (Delta Time): Enter the incremental or delta time limit. (Example: Enter 12:00 hours.) If there are no incremental or delta level components to the prescription, leave these fields (ΔRate and ΔTime) set at their default settings of "0" and press "YES/ENTER" to move to the next field.

d. Max BOLS (Maximum Bolus Dose): Use this field to set the upper limit of the Patient Bolus or Patient Demand dose. If there is no bolus dose, enter "0." If a value is given for the Bolus Dose, use the numeric keys to enter it on this field, press the "YES/ENTER" key to accept, and move to the next field. The value entered for this field must be equal to or less than maximum Bolus Dose listed in Figure 6.1 Administration Route Default Setings table.

e. MinINT (Minimum Bolus Interval): Use this field to set the minimum amount of time allowed between patient bolus doses. When the correct time is entered, press the "YES/ENTER" key to accept, and move to the next field.

f. #BOLS/hr (Number of Boluses allowed per hour): Use this field to enter the maximum patient bolus/hr. When correct, press the "YES/ENTER" key to accept, and move to the next field.

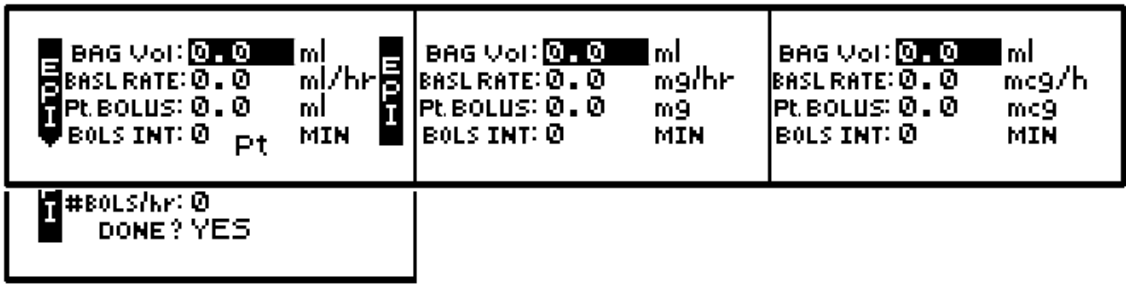
g. ACCEPT? If there are any changes to be made, use the up or down arrows to move to that field, use the numeric keys to enter the corrected data, and press "YES/ENTER" to accept the new values. Move the cursor back to the "ACCEPT?" field and press "YES/ENTER". The medLIMITS are now entered and the display returns to the Pre-Prescription screen. If the entire medLIMITS Setup Screen is to be aborted, press "NO" to return to the Pre-Prescription screen.

6 NEXT? If any field is not satisfactory, use the arrow keys to locate the field to be changed, enter the changes, press the "YES/ENTER" key to accept the new values and then go back to the "NEXT?" field. When all fields are satisfied with correct settings, press the "YES/ENTER" key to exit this screen, and move on to the next PCA Programming Menu.

At the following screen, the patient's prescription information is entered to complete the programming of the pump for a PCA Therapy. Remember that there may be additional limits to rates and doses based on limits for the route of administration selected. (See Figure 6.1).

Note that the screen Identifier gives the abbreviation "EPI" for Epidural(or "PCA" for Patient Controlled Analgesic Therapy if a route other than epidural is selected) and has a down arrow configuration, which indicates that there are more than four fields of text or information to be considered.

EPI Prescription Menu



Note: The PCA therapy can be programmed for Basal Rate (continuous infusion) only, for Bolus (patient demand dose) only, or for both. At least one of these settings must be entered, though, before this menu can be accepted at the "DONE?" prompt.

1 BAG Vol (IV Bag Volume): Using the numeric keys, enter the amount of fluid in the IV Bag or Medication Reservoir (include the overfill volume).

Note: The volume used for priming via the pump is deducted from the Bag Volume. Therefore, Bag Volume should be equal to or greater than the Amount To Be Infused to start the infusion.

When the correct amount has been entered, press the "YES/ENTER" key to accept this field and move on to the next. The range for this field is 1 to 9999.

2 BASL RATE (Basal Rate): The **Curlin Infusion** pump in PCA Epidural Therapy can be set for a continuous basal infusion of 0 to 25 ml per hour. (see Figure 6.1). Use the numeric keys to enter the prescribed rate (the "units" chosen in the Pre-Prescription Menu will appear here).

Note: If the rate falls out of the range of the pump (depending on the administration route selected), a beep will sound, and one of the following alert messages will appear:

This field will not be accepted with the "YES/ENTER" key until the rate is adjusted to fall within pump's range. When the rate is correct, use the "YES/ENTER" key to accept, and move to the next field.

3 Pt. BOLUS (Patient Bolus Dose): Use the numeric keys to enter the prescribed bolus (patient demand dose) and when correct, press the "YES/ENTER" key to accept and move on to the next field. The Patient Bolus Dose can be set for 0 to 25 ml.

Note: If a bolus dose entered falls out of the range of the pump, the above "Greater than Max Field Limit" or "Less than Min Field Limit" alert will also appear and display the limit.

4 BOLS INT (Patient Bolus Interval): This field allows the clinician to enter the minimum amount of time (from 0 to 60 min) prescribed between patient-activated bolus dosing. Enter the time using the numeric keys and, when correct, press the "YES/ENTER" key to accept and move the cursor to the next field.

5 # BOLS/hr (Number of Patient Bolus Doses per hour): Using the numeric keys, enter the number (0 to 15) of boluses allowed per hour and, when correct, press the "YES/ENTER" key to accept and move to the last field.

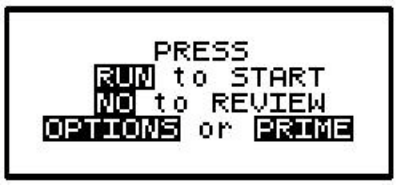
6 DONE?: If the programming is not done, use the up or down arrow keys to locate the field to be changed, use the numeric keys to enter the corrected information and press "YES/ENTER" to accept the new data. If all the fields are acceptable, at the "DONE?" prompt, press the "YES/ENTER" key to move on to the next screen.

IMPORTANT INFORMATION

Before the infusion is started, it is important to consider a few remaining issues, such as checking the administration set to be sure it was primed or purged of all air, or perhaps the pump may need to be put into a security lock setting for additional safety.

The following screen offers you four choices:

Run Options Screen



Note that you do not need to answer yes or no questions or enter data as previously requested. This screen requires only that you press the corresponding key on the keypad to begin the requested function. The following section explains each selection beginning with Prime:

Prime

The IV bag or syringe and administration set should be inspected and be free of all air and air bubbles before they are connected to the access site. Priming may have been done by gravity when the administration set was first connected to the bag (see Chapter 2). However, if it was not done then or if there are any remaining air bubbles to be removed, the pump's "PRIME" function will assist in this process. To prime the set by using the pump, follow the directions below.

Note: The lock setting may restrict the ability to prime using the pump.

1 Press and release the "PRIME" key at the "Run Options" screen.

2 The following screen appears:

Prime Direction Screen

3 Disconnect the administration set from the patient's access site using aseptic protocols.

IMPORTANT INFORMATION

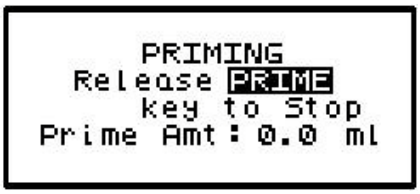
Priming with the set connected to the patient could result in overdose and may cause injury or even death to the patient.

4 Release any clamps on the administration set.

5 Press and hold the "PRIME" key. Fluid will be pumped at a rate of 300 ml/hr through the administration set as long as the PRIME key is pressed or until 6 ml is pumped. The pump will then beep once and require repressing and holding the "PRIME" key again until 6 more milliliters is delivered. Continue this process until all air has been purged from the IV bag and the administration set.

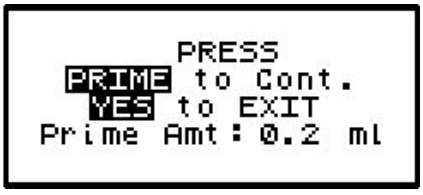
Note: If the bag volume is depleted during priming, a subsequent priming event cannot be initiated.

While the "PRIME" key is depressed, the following screen appears:



The amount used for the prime session is accumulated and displayed on the bottom field of this screen.

6 When priming is complete, release the "PRIME" key. The following screen will display:



7 To continue priming, press the PRIME key again. If priming is complete, press "YES/ENTER" to exit and return to the Run Option screen. The total amount of solution used in this priming session displays on the bottom field.

Note: The amount of fluid used in the priming process will be deducted from the Bag Volume Total but will NOT be counted in the amount infused because this amount is not to be

delivered to the patient. The bag volume should still be equal to or greater than the Amount TBI before you can proceed with the infusion. If it is not, an alert will appear stating, "AMT TBI GREATER THAN BAG VOL." You have the option to continue by pressing the "YES/ENTER" key or return to the "Run Options" screen by pressing the "NO" key.

Options

Once the prescription is programmed, there are more customizing features available for each therapy. Press the "OPTIONS" key from the Run Options screen to access these options. The OPTIONS menu can also be accessed while the pump is running.

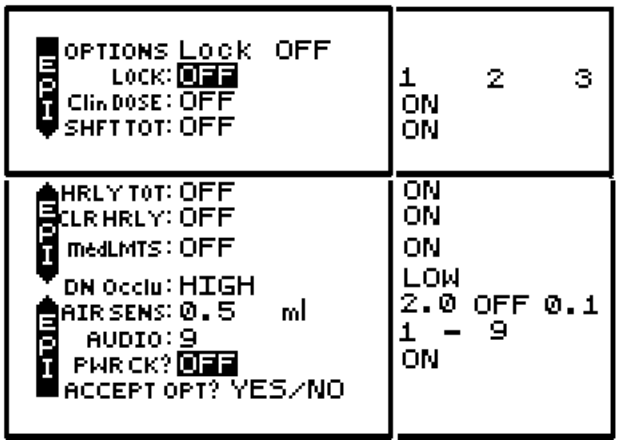
Note. After the infusion has started, any changes made through the OPTIONS menu involving:

- "Clin DOSE",
- "SHFT TOT"
- "HRLY TOT", and
- "CLR HRLY"

will return you directly to the "ACCEPT OPT?" field.

The following screen displays the EPI Therapy "Options" available. When you open this menu, use the up and down arrows to move the highlight bar to the desired option. Lock settings may require an access code before the option can be changed. Any new setting, except "CLEAR HOURLY" screen acceptance and "SHIFT TOTAL" clear screen acceptance becomes effective when the "Options" screen is exited by pressing "YES/ENTER" at the "ACCEPT OPT?" field.

EPI Options Menu

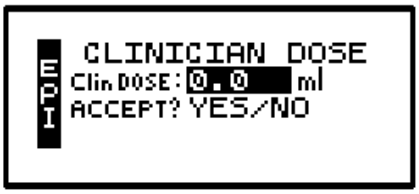


Note: The pump's default settings are indicated above in the **left** screen.

1 LOCK: When the lock is set to "OFF," the clinician has access to all of the pump functions. After the pump is completely programmed, change the lock to meet the security needs of this therapy. There are four choices: OFF, 1, 2, or 3. Each level higher limits the available options. To change the lock setting, press the "NO/CHANGE" key until the correct lock setting appears. Then, press "YES/ENTER" key to accept the new lock selection. Once the pump has been set into a locked condition, any subsequent request to change the lock will require a clinician access code. (See "Clinical Information and Access Code" chapter for access code information, lock setting table and available features for each lock setting).

2 ClinDOSE? (Clinician Dose): Allows an additional prescribed dose to be administered by a clinician at any time except during a Load Dose or Bolus Dose. The Maximum Clinician Dose is determined by the administration route (see Figure 6.1).

Clinician Dose Menu



Note: If a lock setting of 1, 2, or 3 is selected, an access code will be requested. If you wish to give a clinician dose at this time, change OFF to ON and press YES key. Enter the amount of clinician dose to be delivered. Press YES to accept the clinician dose menu. The Clinician Dose will begin when the options screen is accepted and the pump is running. The following Run Screen will appear until it is completed:

Clinician Dose Run Screen

The graphic bar now depicts the clinician dose in progress and when this dose is complete, reverts back to reflecting the container in progress. **The graphic bar will toggle with Line Pressure ("L Press") which displays the current patient side (downstream) pressure in the administration set in millimeter of mercury (mmHg) units.**

The Volume Infused will toggle with Dose Infused if mg or mcg are the selected unit. Remaining dose time will toggle with remaining dose volume to infuse.

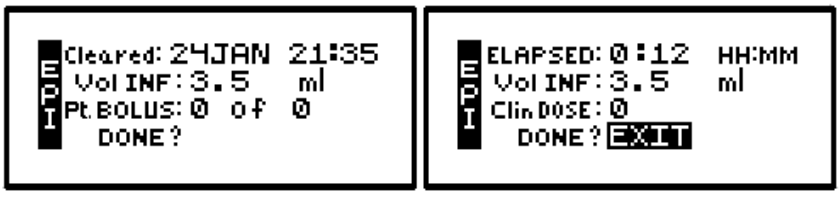
IMPORTANT INFORMATION

When the clinician dose is complete, the pump will beep, return to the previous running rate, and resume the PCA therapy. <u>The lockout time for bolus dosing will restart at the end of the clinician dose to prevent a bolus from being delivered too quickly after a clinician dose</u>.

Once the clinician dose is given, the clinician-dose field returns to its default of "0," and any subsequent clinician doses will need to be reentered.

3 SHIFT TOT (Shift Totals): This feature allows the clinician to track the infusion for a selected period of time, such as an 8 or 12-hour shift. To access Shift Totals, press the "NO/CHANGE" key to change OFF to ON and then press "YES/ENTER." The following screens will appear alternately:

Shift Totals Screens



a. Cleared/ELAPSED: This field displays the date and time the shift total was last cleared. It toggles with the amount of time elapsed since that clearing.

b. Vol INF/Amt INF (Volume/Amount Infused): This field displays the volume (ml) and amount (mg or mcg) infused since the shift total was cleared.

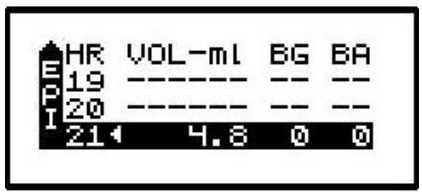
c. BOLUS/Clin DOSE (Clinician Dose): This field displays the number of clinician doses given since the total was cleared. It toggles with the number of patient activated boluses received of those requested.

d. To exit the shift total feature without clearing the totals, press the "YES" key at the highlighted EXIT field.

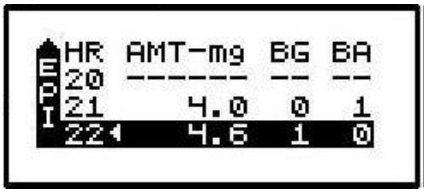
e. To clear the Shift Totals, press the "NO" key at the EXIT field. The field will change to read "CLEAR", press the "YES" key now to clear. A lock level 3 will require an access code entry before allowing the user to clear the Shift Totals.

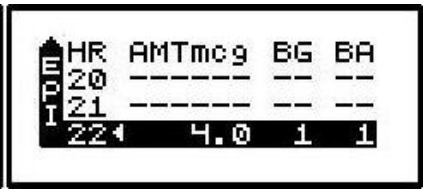
4 HRLY TOT (Hourly Totals): To view hourly infusion totals press the "NO/CHANGE" key to change OFF to ON and then press "YES/ENTER". Either of the following two screens will appear:

Hourly Totals Screen



or





The screen displays hourly infusion delivery for each hour (00-23) of the last 24 hours of therapy in the unit selected.

a. The highlighted field has a marker ◀ indicating the current hour in progress.

b. Volume or Amount will show in the first column, BG (Patient Bolus Given) in the second column and BA (Patient Bolus Attempted) in the third column.

c. Use the up arrow key to scroll back through the last 24 hours.

d. If the infusion was not running, a series of ------ will appear for that hour.

e. To exit this screen, press the NO/CHANGE key.

5 CLR HRLY (Clear Hourly Totals): To clear Hourly Totals press the "NO/CHANGE" key to change OFF to ON and then press "YES/ENTER." The following screen will appear:

Clear Hourly Totals Screen

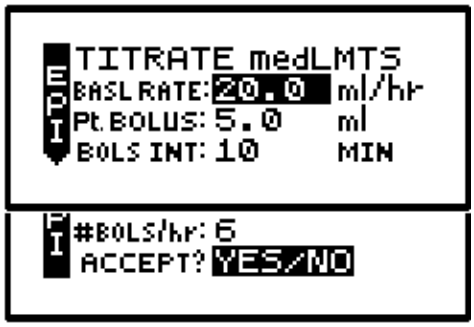
a. To clear the Hourly Totals, press the "YES/ENTER" key. All Hourly Totals will be cleared and the cursor will return to the Options menu.

b. To maintain the Hourly Totals, press the "NO/CHANGE" key. Hourly Totals will be maintained and the cursor will return to the Options menu.

c. A lock level of 2 or 3 will require an access code entry before allowing "Hourly Totals" reset.

6 medLMTS™ (Medication Limits): If this feature was **not** selected in the Pre-Prescription Menu, it will not appear on the Options menu. If medLIMITS™ parameters were programmed previously, this field will appear; and if you want to titrate now, select "ON" and press the "YES/ENTER" key to go to the PCA Titrate medLIMITS™ menu:

EPI Titrate MedLIMITS™ Screen



This screen allows you to change the basal rate and/or the bolus parameters within the predefined prescription limits. Whenever any value is changed using the medLIMITS™ option, the new value will appear on the prescription menu when it is reviewed. If a delta time/rate limitation was previously programmed, the rate can be increased within those parameters up to 12 times during the delta time period. If a 13th attempt is made an alert will display: "MAX CHANGES FOR PERIOD REACHED."

a. BASL RATE: Use the numeric keys to enter the new desired rate. When the correct rate has been entered, press the "YES/ENTER" key to accept the rate, and move to the next field. The range for this field is 0 to 25.

b. Pt. BOLUS: Use the numeric keys to enter the new patient bolus dose and, when correct, press "YES/ENTER" to accept and move to the next field. The range for this field is 0 to 25.

c. BOLS INT (Patient Bolus Interval): Use the numeric keys to enter the new bolus time interval and, when correct, press "YES/ENTER" to accept and move to next field. The range for this field is 0 to 60.

d. #BOLS /hr (number of patient boluses per hour): Use the numeric keys to enter the new number of patient boluses per hour and, when correct, press "YES/ENTER" to move to the final step in the medLIMITS Setup menu. The range for this field is 0 to 15.

e. ACCEPT? YES/NO: If necessary, use the arrow keys to move to any field and change the data entered. To abort this screen entirely, press "NO" at this prompt. Any data entered on this display will be aborted and the display will return to the options menu . If the parameters entered are correct, press the "YES/ENTER" key, and the new rate or rates will be accepted. The display returns to the Options menu when this screen is exited.

f. New values will be in effect when the Options menu is accepted.

7 DN Occlu (Down Occlusion): This feature allows the pump to sense an occlusion pressure in the line between the pump and the access site, e.g., a kink in the tubing between the pump

and the patient, a blockage in the access site, or a clamp left on the distal tubing. There are two options for this field: the "Low" setting triggers the alarm when it detects in-line pressures at 400 mmHg ±150 (8 ±3 psi "pounds per square inch"), or the "High" setting (default setting) alarms when it detects in-line pressures at 900 mmHg ±150 (18 ±3 psi). Press "NO" to change the selection. When the correct pressure setting appears, press the "YES/ENTER" key to accept it and move the cursor to the next field.

8 AIR SENS (Air In Line Sensor): This feature allows you to select the amount of air the pump will detect in the administration set tubing before it goes into an alarm state. The choices are 0.1 ml, 0.5 ml (the default setting), 2.0 ml, and OFF. Press the "NO/CHANGE" key to change the options until the desired setting appears on the display, and then press "YES/ENTER" to select and move the cursor to the next line.

IMPORTANT INFORMATION

Whenever the Air-In-Line Sensor is set to OFF, Moog recommends the use of an air-eliminating filter.

When the Air-In-Line Sensor is set to "OFF" and the "RUN" key is pressed at the Run Option screen, the alert message below will appear on the screen. It will also appear if the Air-In-Line is changed from the OPTIONS menu when an infusion is in progress and the AIR SENS field is accepted.

Air-In-Line Sensor Off Message Screen

This message alerts you to the Air-In-Line Sensor being set to "OFF." The message will require answering "yes" or "no" to whether the administration set being used has an air-eliminating filter. If you press the "YES" key to indicate that there is an air-eliminating filter in use, the Air-In-Line sensor will be completely disabled and the pump will start the infusion.

If you press the "NO" key to indicate a filter is not in use, the Air-In-Line sensor will be set to detect the presence of any contiguous 2-milliliter air in the line. The alert message below will then briefly appear on the screen. If the alert occurred at the Run Options screen, the pump will start the infusion.

Pump Will Sense 2ml Alert Screen

Also, when the Air-In-Line sensor is completely disabled and the infusion goes to an "Infusion Complete" or "Empty Bag" status, the Air Sensor setting will automatically change to 2.0 ml.

9 AUDIO (Audio Volume): This feature allows you to adjust the audio volume of the pump's alarm setting. You can enter a number from 9 (the loudest setting) to 1 (the quietest setting). When you select a number, the pump will beep at that audio level. Press the "NO" key and

enter a new setting (1 – 9). Repeat until the desired setting is reached, and then press "YES/ENTER" to select and move the cursor to the next field.

10 PWR CK? (Power Check): This feature allows you to periodically recheck the battery charge. It will display the active power source of the pump, whether internal batteries or external power. If the active source is the batteries, a graph will indicate approximately how much charge remains in the batteries. To access this information, select "ON" and press "YES/ENTER". The Power Source Screen will display for five seconds and return to the Options Menu.

11 ACCEPT OPT? If the changes made in the Options Menu are acceptable, press the "YES/ENTER" key. If the changes are not acceptable, cursor up to any field for corrections or abort all changes made in this menu by pressing the "NO" key, and return to the previous screen.

There are two remaining selections to consider from the "Run Options Screen", **NO** to **REVIEW** and **RUN** to **START**.

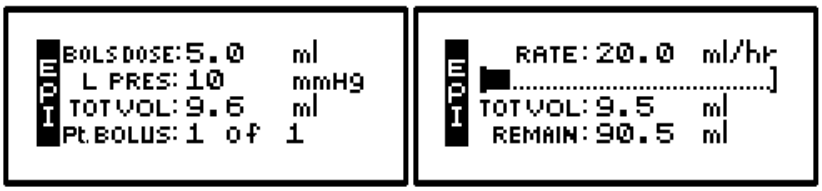
Reviewing or Changing the Program

To review or make any changes in the prescription programming, press the "NO/CHANGE" key at the Run Options screen, and the prescription screens will reappear (Changes may be inhibited by lock setting). Changes to the prescription can be made before the start of the therapy; however, once a therapy has started, no changes can be made to the Pre-Prescription Screen. It will appear only for review purposes. Attempts to change the Pre-Prescription Menu during a new program review will cause an alert screen to be displayed, notifying the programmer that changes will require reprogramming of the prescription.

Starting the Infusion

RUN: When the programming is complete and the pump has been primed properly and connected to the access site, press the "RUN" key at the Run Option screen to start the therapy. The pump will begin infusing, and except when a LOAD DOSE or CLIN DOSE is programmed, the following screen will display data on the status of the therapy:

EPI Run Screen



IMPORTANT INFORMATION

The amount of drug infused during the "loading dose" depends on the units chosen which may be in ml's, mg, or mcg. A loading dose of 5 mg will infuse 50 ml if the concentration is 0.1 mg/ml.

This screen will continue to display as long as the infusion is running. Notice the information on each field:

1 The first field displays the "RATE" of the infusion toggled with the programmed Patient

Bolus Dose Amount. This field also displays "BOLUS xx (mg/ml/mcg)" whenever a bolus dose is being delivered. The programmed bolus amount displays and the BOLUS text flashes when the pump is delivering a patient bolus dose.

2 The second field is a graphic bar representing the percentage of the completed infusion. **The graphic bar will toggle with Line Pressure ("L Press") which displays the current patient side (downstream) pressure in the administration set in millimeter of mercury (mmHg) units.**

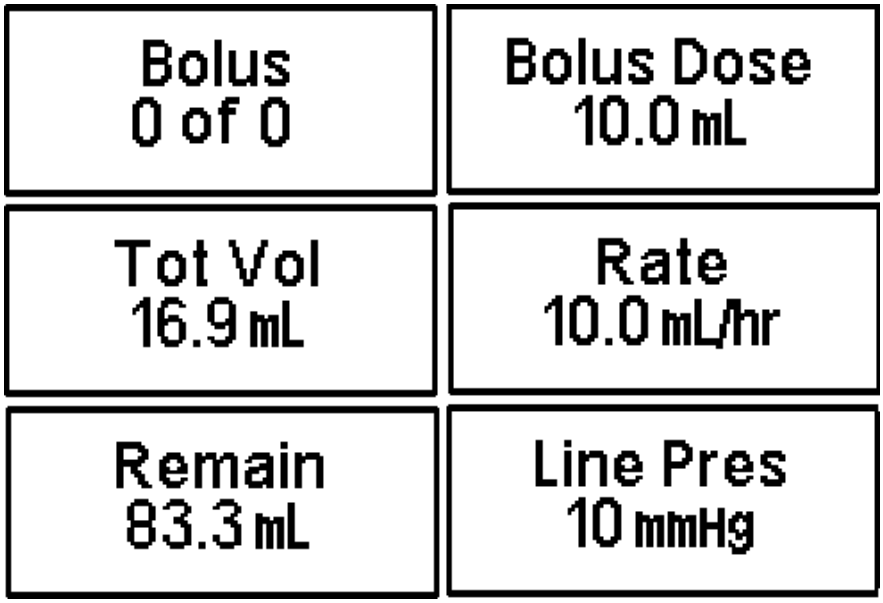
3 The third line displays the total volume "TOT Vol" infused and, if mg or mcg is the selected unit, this field will toggle to display the total mg or mcg infused "TOT INF" as well.

4 The fourth line displays the number of patient boluses delivered and number of boluses attempted per bag volume. If the patient attempts a bolus before the allowed amount of time set between boluses, an alert message will display "Bolus Requested," and only the second number will increase. Also, toggling with number of boluses delivered/requested is the remaining volume left in the bag.

5 In addition to the display screen, information on the status of the infusion is provided by the three LED lights. The green LED light blinks whenever the pump is infusing. If the Basal rate is set at "0," the yellow LED light will blink along with the green light until the bolus is actually being delivered. While the bolus is infusing, only the green light will blink.

The information on the run screen will alternate between a four-line display and a large font display of each individual line of information on the screen.

EPI Information Screens



Interrupting an Infusion

The infusion can be paused at any time by pressing the "PAUSE" key. The yellow "Standby" LED light blinks, and the following screen displays:

Pause Menu Screen

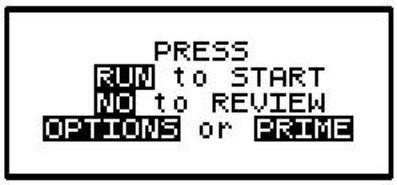
The screen offers the user three options:

Resuming the Current Container

1 RESUME: To resume the therapy in progress from where it left off, press the "YES/ENTER" key at this field. The pump will return to the Run Options Screen, and you can resume the therapy.

Note: If you wish to hang a new container, move the cursor to REPEAT Rx (see following section).

Run Options Screen



a. Press the "RUN" key to resume the therapy.

b. Press the "NO" key to go back to the programming screen to make changes in the therapy program (this feature may be blocked by a security lock, thus not allowing changes to the program unless an access code is given). Also, remember that no changes are allowed in the first prescription menu if an infusion is in progress.

c. Press the "OPTIONS" or "PRIME" keys if either of these functions is desired at this time.

Hanging a New Container

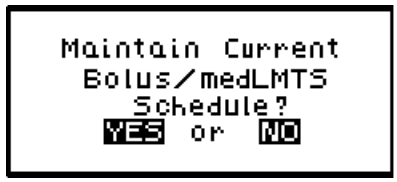
2 REPEAT Rx: To repeat the present therapy and add another IV container with the same prescription, press the "YES/ENTER" key at this field. If more than one therapy is programmed, a "Select Therapy Screen" will list all preprogrammed therapies. Select which therapy to repeat by moving the cursor to the therapy of choice and pressing the "YES/ENTER" key.

The pump will then display the program screen with the last programmed values of the therapy and the cursor will automatically advance through the prescription menus. You can stop the auto-advance function by pressing any key (except ON/OFF) and then complete the confirmation of the prescription manually by pressing the "YES" key at the remaining fields.

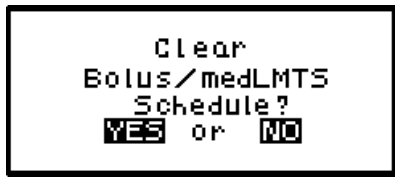
When the program is accepted, the display returns to the above "Run Options Screen." If the current therapy is still in progress when the "REPEAT Rx" function is selected, the following screen will appear:

a. This screen is a reminder that the infusion is not complete. If the incomplete therapy is to be continued, press "NO/CHANGE" and the Pause Menu screen will be displayed, then press "YES/ENTER" and the Run Options screen will be displayed. To resume the unfinished therapy, press "RUN."

b. If the therapy is to be terminated and a new container added, press "YES/ENTER" to repeat the infusion. The following screen will then appear, asking whether or not you wish to maintain the current Bolus or MedLIMITS schedule.



Press "YES/ENTER" to maintain the current schedules and continue with the programming of the pump. Press "NO/CHANGE" to start a new schedule. The following screen will then appear asking you to confirm the clearing of the schedules.



Press "YES/ENTER" to clear the current schedules and continue with the programming of the pump. Pressing "NO/CHANGE" will return you to the prior screen.

Entering a New Program

3 NEW PROGRAM: To program an entirely new prescription or different therapy, press the "YES/ENTER" key at this field (the ability to change these parameters may be controlled by the lock setting). The display will return to the "Pre-Prescription Screen." If the current container has not completely infused, the following screen will appear:

a. This screen is a reminder that the infusion is not complete. If you want to return to the current container and finish infusing it, press the "NO/CHANGE" key to return to the Pause Menu screen.

b. To abort the current container and enter a new program, press the "YES/ENTER" key. If a prescription has previously been entered for the therapy mode you wish to program, the following screen will appear:

c. If you want to save the current prescription, press the "NO/CHANGE" key to return to the Pause Menu screen. If you wish to erase the current prescription and enter a new program, press the "YES/ENTER" key to proceed. All previous prescription values will be erased.

Note: Selecting "NEW PROGRAM" will also reset the OPTIONS selected to their default settings except for the LOCK and AUDIO settings which must be changed manually.

Infusion Complete

When the infusion is complete, the audio alarm will beep, and the following screen will display:

Note: To temporarily silence the alarm, press the "Silence" key.

PCA Infusion Complete

This screen displays:

1 "INFUSION COMPLETE."

2 A completely filled in bar graph indicating infusion complete.

3 TOT Vol (Total Volume): If units are mg or mcg, this field will also toggle to give the Total Infused.

4 Infusion Complete is considered an alarm status, and the audio alarm will beep periodically every three seconds to alert the user to this condition. Additionally, the red LED light blinks to indicate that the "Infusion Complete" alarm exists.

Stopping the Infusion

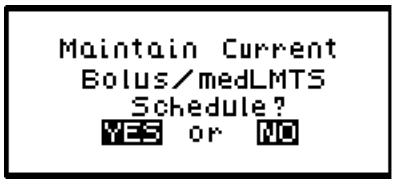
At "Infusion Complete", press the "PAUSE" key and then choose to either:

1 Turn off the pump by pressing the "OFF" key, or

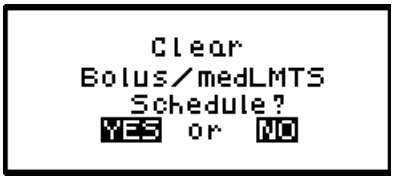
2 Select one of the options from the "Pause" screen:

Pause Menu Screen

a. REPEAT Rx: To repeat the present therapy and add another container with the same prescription, press the "YES/ENTER" key at this field. The following screen will then appear, asking whether or not you wish to maintain the current Bolus or MedLIMITS schedule.



Press "YES/ENTER" to maintain the current schedules and continue with the programming of the pump. Press "NO/CHANGE" to start a new schedule. The following screen will then appear asking you to confirm the clearing of the schedules.



Press "YES/ENTER" to clear the current schedules and continue with the programming of the pump. Pressing "NO/CHANGE" will return you to the prior screen.

The pump will then display the program screens with the last programmed values of the therapy and the cursor will automatically advance through the prescription menus. You can stop the auto-advance function by pressing any key (except ON/OFF) and then complete the confirmation of the prescription manually by pressing the "YES" key at the remaining fields.

b. NEW PROGRAM: To program an entirely new prescription press the "YES/ENTER" key when this field is highlighted and the display will return to the "Select Therapy Screen." (The lock setting can control access to this function.)

Note: The therapy is complete; therefore, RESUME is not an option at this time and is not displayed.

Chapter 7

Troubleshooting

This section assists in identifying and resolving alert and alarm conditions that may occur during the use of the ***Curlin Infusion*** *pump.*

Alerts, Alarms, and Error Codes

IMPORTANT INFORMATION

Alerts, Alarms, and Error Codes are classes of events that are relayed to the user and logged into the history file. The screen will display messages, the LED status lights will blink, and the audio alarm will sound. The backlight feature will illuminate the screen during an alert or alarm.

If the batteries have been dislodged or are so low in power that they cannot operate the pump, a solid tone may occur accompanied by a blank display screen and a constant red alarm light. In this event, replace the batteries.

ALERTS

Alerts report conditions that require attention from the user but may not be safety issues. When an alert occurs, a message is displayed on the LCD screen, and the audio alarm is sounded. The message displays on the screen for three seconds if it is a user-initiated alert, such as a wrong key press. If it is an automatic alert occurring while a therapy is infusing, such as "low battery," the message displays for twenty seconds. If it is an alert that requires a user response, the alert remains displayed until the user presses one of the keys specified on the screen. The following is a list of the alerts in alphabetical order.

LCD Display: ALERT AIR SENS OFF. Using In Line Filter?

Alert: Air-In-Line Sensor Disabled

Cause: The Air-In-Line Sensor has been disabled and either the "RUN" key has been pressed to run an infusion or the AIR SENS setting has been set to OFF on the Options menu while the pump is running.

Resolution: If an air-eliminating filter is being used, press YES. The infusion will start/resume and the Air In Line sensitivity will be disabled until the empty bag alert occurs or infusion complete is reached. No "AIR IN LINE" alarms will occur while the air sensitivity is disabled.

If no air-eliminating filter is being used, press NO. The infusion will

start/resume and the Air In Line sensitivity will be set to 2 ml for the remainder of the infusion.

Note: If the screen is not acknowledged with a YES or NO key press, the Air In Line sensitivity will be set to 2 ml when the screen is cleared (i.e. alarm occurs while screen is displayed).

LCD Display: ALERT AIR SENSOR HAS BEEN SET TO 2.0 ML

Alert: Air in line sensitivity will be set to 2.0ml

Cause: NO was pressed at the "AIR SENS OFF. Using In Line Filter?" alert screen. The air in line sensitivity will be set to trigger an AIR IN LINE alarm if 2.0ml of air is detected.

Resolution: If an air eliminating filter is being used, press YES when the "AIR SENS OFF. Using In Line Filter?" screen is displayed. This will disable air detection as indicated under "AIR SENS OFF. Using In Line Filter?" alert explanation.

LCD Display: ALERT AMT NOT 0 OR CURRENT DOSE AMT

Alert: Unable to change clinician dose while clinician dose in progress

Cause: A clinician dose is in progress, and a new clinician dose value is entered that is not "0" or the current dose amount.

Resolution: If the current clinician dose is to be continued, change the 'Clin DOSE' value back to the original value and accept the field and menu.

If the current clinician dose is to be terminated, which will allow a new clinician dose to be programmed, enter a value of 0 in the 'Clin DOSE' field and accept the field and menu. The next time that the Options menu is accessed a new 'Clin DOSE' value can be entered.

LCD Display: ALERT AMT TBI GREATER THAN BAG VOLUME

Alert: Value entered is greater than value in Bag Vol field.

Cause: Occurs when the amount the entered for the "Amt TBI" or "Vol TBI" field is greater than the value in the bag volume field.

Resolution: Recheck the data entry.

LCD Display: ALERT AMT TBI GREATER THAN BAG VOLUME. YES to Continue NO to Return

Alert: Insufficient volume

Cause: Occurs when bag volume is less than amount to be infused. Can also occur when pump's priming feature is used and bag volume is depleted by the amount used in priming.

Resolution: To modify the Bag Volume or Amt/Vol TBI, press "NO" to return to the Run Options screen. The programming screens will then be accessible by pressing "NO" to Review on the Run Options screen.

Note: Access to the programming screens is lock level restrictive.

If the actual amount in the source bag/container is sufficient to complete the programmed infusion, press "YES" to start the infusion.

Note: If the infusion is continued without sufficient volume, air-in-line or up occlusion alarms may occur later in the infusion.

LCD Display: ALERT AMT TBI LESS THAN AMT INFUSED

Alert: Unable to change amount to be infused to an amount less than what has already infused.

Cause: Occurs when user tries to change the Vol TBI / Amt TBI field in the prescription menu to an amount lower that the amount which has already infused.

Resolution: Recheck the data entry. If alert recurs, resume the infusion and observe the Vol Inf / Amt Inf field on the run menu. The value for data entry must be greater than the value displayed on the run menu.

LCD Display: ALERT BOLUS DOSE IS NON-ZERO

Alert: Unable to enter a bolus interval or # of boluses/hour of "0"

Cause: Bolus interval or # of boluses/hour value entered is zero, and bolus dose value is non-zero. When a bolus value is entered, there must also be a value greater than 0 entered for bolus interval and # of boluses/hour.

Resolution: Recheck data entered for the prescription.

LCD Display: ALERT BOLUS DOSE NOT ALLOWED

Alert: A bolus dose is not allowed

Cause: Bolus key or button pressed when no bolus dose is programmed.

Resolution: If bolus doses are part of the prescription, modify the Rx/medLIMITS program to allow bolus doses.

LCD Display: ALERT BOLUS REQUESTED

Alert: Not time for bolus

Cause: Bolus button or key is pressed before the programmed time interval between bolus doses have elapsed.

Resolution: Advise patient of time limits between bolus doses per agency protocols.

LCD Display: ALERT CALCULATED AMT OUT OF RANGE

Alert: Invalid calculated amount for infusion

Cause: The calculated amount for the infusion, based on the values entered for Rate and Time, exceeds either the pump limit or the value entered for bag volume.

Resolution: Recheck data entry for Rx.

LCD Display: ALERT CALCULATED RATE OUT OF RANGE

Alert: Invalid calculated rate of infusion

Cause: The calculated rate for the infusion, based on the values entered for Volume/Amount and Time, exceeds either the minimum or maximum rate limit of the pump or the maximum rate programmed in medLIMITS.

Resolution: Recheck data entry for Rx.

LCD Display: ALERT CALCULATED TIME OUT OF RANGE

Alert: Invalid calculated time

Cause: The calculated time for the therapy, based on the values entered for Volume/Amount and Rate, exceeds the therapy time limit of 288:00 HHMM (167:59 HHMM Software Versions below R37C).

Resolution: Recheck data entry. The time calculated by the pump cannot exceed the pump range.

LCD Display: ALERT CHANGES REQUIRE REPROGRAMMING. YES to Continue NO to Return

Alert: Changes to the data will require re-entry of prescription.

Cause: Attempted to change the Pre-Prescription Screen prior to start of infusion.

Resolution: If a change to the programmed prescription is required, press "YES".

Note: Selecting "YES" will clear the entire prescription and require re-entry the prescription.

If no change is required, press "NO" to preserve the current prescription and return back to the Pre-Prescription screen.

LCD Display: ALERT CMS LEARN MODE ACTIVE – KEY NOT ACCEPTED

Alert: The pump is performing a CMS Create Prescription function and the key pressed is not valid for this mode.

Cause: While using the pump with CMS/PC in order to create a prescription only certain keys are allowed to be recorded as part of the prescription. Whenever a prohibited key is pressed (i.e., RUN key) this alert will occur and the key will not be recorded.

Resolution: Do not press the key causing the alert during a CMS Create Prescription function.

LCD Display: ALERT DATE/TIME CHANGE NOT ALLOWED

Alert: Not allowed to change date or time

Cause: Attempt made to modify system clock while a therapy/delay is in progress.

Resolution: If the therapy in progress is to be continued, wait until the infusion is complete and then change the Date and Time.

If it is not necessary to preserve the therapy in progress or any other programmed therapy, use the "CLEAR Rx" feature of the Clinician Biomed menu. This feature will erase all programmed therapies in the pump. The Date and Time can then be changed.

If the therapy is to be terminated but the prescription preserved (excluding Delayed Start), go to the "Pause" menu, select "REPEAT Rx" and press "YES" at the "Rx in progress" alert screen. The pump can then be turned off and then on to access the Date/Time change option in the Clinician Biomed menu. Selecting "NEW PROGRAM" and the name for the therapy in progress, instead of "REPEAT Rx", may also be used but this selection will erase the prescription for the therapy.

LCD Display: ALERT DATE/TIME INVALID

Alert: Unable to enter invalid date or time

Cause: Date or time entry is not in a valid format.

Resolution: Recheck data entry. An invalid date or time will not be accepted.

LCD Display: ALERT DELAY START TIME NOT VALID

Alert: Delay Start time not valid

Cause: Delay start time entered is less than one minute or greater than 24 hours from the current system time.

Resolution: Reset Delay Start Time to fall within pump limits. (To obtain the current date and time of the pump, turn the pump off and then on and observe the time on the Welcome screen).

LCD Display: ALERT DOOR OPEN or SET NOT PROPERLY INSTALLED

Alert: Door Open or Administration Set is not properly installed

Cause: Door Latch is not completely closed or the Administration Set is not installed according to installation instructions.

Resolution: Recheck the door latch assembly and securely close the door. If the door was closed properly, re-install the set according to the installation instructions.

LCD Display: ALERT DOSE AMT IS LESS THAN DOSE INFUSED

Alert: Dose amount entered is less than the amount already infused for the dose.

Cause: New dose value entered is less than the amount already infused for the dose.

Resolution: Recheck data entry. After the dose in progress finishes, the dose amount may be changed to the lower value.

If the above is not suitable, re-program the prescription.

LCD Display: ALERT DOSE CANCELLED, RESUME INFUSION, THEN NEW DOSE

Alert: Cancellation of a Clinician Dose is pending and can only be processed with the pump running.

Cause: Clinician Dose in progress is cancelled while the infusion is paused; and the user attempts to enter a new Clinician Dose, prior to resuming the infusion

Resolution: Resume the infusion and then program the desired Clinician Dose.

LCD Display: ALERT DOSE FREQ LESS THAN DOSE TIME

Alert: Dose frequency time is less than the time required to infuse the programmed dose

Cause: The dose frequency value entered is less than the time required to infuse the programmed dose amount.

Resolution: Recheck data entry. Dose frequency cannot be less than required time for dose to infuse.

LCD Display: ALERT EMPTY BAG

Alert: Empty Bag

Cause: Occurs when the volume infused for the prescription is equal to the "Bag Vol". Bag Vol may have been reduced due to priming.

Resolution: Verify that the volume remaining in the source container is sufficient to complete the current therapy. If the volume is insufficient, add another source container or stop the infusion.

LCD Display: ALERT EXCEEDS POSSIBLE DOSES PER HR

Alert: Unable to enter # of boluses/hr that would be possible given the programmed bolus amount

Cause: # Boluses/hr value exceeds the possible # given the bolus amount.

Resolution: Recheck data entered. The # of boluses per hour cannot exceed the time required to deliver the bolus amount programmed.

LCD Display: ALERT EXTERNAL POWER DISCONNECTED

Alert: External power source disconnected

Cause: External power source is disconnected or external battery pack power is completely depleted during pump operation.

Resolution: Reestablish external power source, or let pump continue to operate on batteries.

Note: The battery level should be checked after the external source is disconnected to gage if additional batteries will be required before an external power source is available again.

LCD Display: ALERT FIELD NOT ACCEPTED

Alert: Field Not Accepted

Cause: The field indicated by the flashing cursor has not been accepted.

Resolution: Recheck the field and accept it by pressing the "YES/ENTER" key.

LCD Display: ALERT GREATER THAN MAX FIELD LIMIT, MAX = _____

Alert: Unable to program a field with a value that exceeds the maximum limit

Cause: Value entered is greater than the allowable limit.

Resolution: Recheck data entry. A value entered cannot exceed the field limit.

LCD Display: ALERT GREATER THAN WHAT BOLS INT ALLOWS

Alert: Unable to enter a # of boluses/hr that exceeds the possible number allowed given the bolus interval

Cause: # Boluses/hr value exceeds the possible number given the bolus interval ("BOLS INT") programmed.

Resolution: Recheck data entry. The # of boluses/hr cannot be set for higher than the number allowed using the given bolus interval.

LCD Display: ALERT INFUSION NOT IN PROGRESS

Alert: Infusion not in progress

Cause: Modification attempted to Next Dose or Early Down Ramp field without an infusion in progress.

Resolution: Wait until infusion is in progress before modifying or modify original Rx programmed.

LCD Display: ALERT INSUFFICIENT VOLUME MAX DOSES=_____

Alert: Bag Volume value is insufficient to deliver the remaining

Cause: Occurs whenever the prescription is changed so that Bag Volume is now less than Amount Required for the entire therapy (based on volume already infused plus volume required to deliver the remaining doses).

Resolution: Recheck data entry. If the source container contains sufficient volume to complete the remaining dose cycles (dose volume + KVO volume), increase the "Bag Vol". If the volume is insufficient, decrease the "DOSE/BAG" such that the therapy can continue with the new modified prescription.

Note: If either "Bag Vol" or "DOSE/BAG" is modified and the therapy is repeated, these fields will have to be verified against the source container volume and prescribed prescription and changed accordingly.

LCD Display: ALERT INVALID ENTRY

Alert: Invalid Entry

Cause: Data entered does not fall within the defined limits of the program.

Resolution: Enter data that falls within the limits of the program. Refer to the specific therapy section in the User's Manual.

LCD Display: ALERT KVO RATE IS GREATER THAN INF RATE

Alert: Unable to enter KVO rate that is greater than infusion rate

Cause: Occurs when user enters a KVO rate that is greater than infusion rate.

Resolution: Recheck data entry. KVO rate must be less than programmed or calculated infusion rate.

LCD Display: ALERT LESS THAN MIN FIELD LIMIT, MIN = _____

Alert: Unable to program a field with a value that is less than the minimum limit

Cause: Value entered is less than the minimum limit for the pump or medLIMITS programmed.

Resolution: Recheck data entry. A value entered cannot be less than the field's minimum limit.

LCD Display: ALERT "LOW C BATTERIES" **Alert:** Battery source power is low

Cause: The batteries that are powering the pump have reached levels that may prevent the continued operation of the pump beyond 1 hour (actual time is rate dependent). Alert applies only to "C size" batteries, not to the rechargeable battery pack.

Resolution: "PAUSE" the therapy in progress, turn the pump "OFF"; and then replace the batteries. (See chapter 1 of the User's Manual for instructions on battery change.)

LCD Display: ALERT MAX CHANGES FOR PERIOD REACHED

Alert: The maximum number of changes for the current delta time window has been reached

Cause: The maximum number of pump rate changes for the current delta time window has been reached. This occurs when the rate has been changed more than 12 times during any delta time window.

Resolution: Allow more time to elapse before attempting to change the rate or reprogram the therapy with medLIMITS and prescription basal and bolus ranges that will reduce the number of times that the rate will have to be adjusted.

LCD Display: ALERT MODIFICATION NOT ALLOWED

Alert: Modification Not Allowed

Cause: The selection cannot be modified at this time or the programmed lock setting does not permit modifications to this selection.

Resolution: Review the User's Manual for program restrictions or if lock level restricted, change lock setting to access area you wish to modify.

LCD Display: ALERT NEW DOSE IS LESS THAN CURRENT DOSE

Alert: Unable to program a lower number of doses than number of doses already infused

Cause: While the Variable or Intermittent therapy is in progress, a lower number of doses to be delivered is entered than the number already infused.

Resolution: Wait until current therapy in progress is complete, or "PAUSE" pump and, using the REPEAT Rx function, modify the number of doses.

Note: Selecting to repeat the therapy in progress will terminate the current therapy.

LCD Display: ALERT NO BOLUS DOSE DEFINED

Alert: Unable to enter bolus interval or # of bolus/hr because no bolus value has been programmed

Cause: Entry of a value other than zero for the Bolus Interval or #Bolus per hour field when no bolus value has been programmed.

Resolution: Enter bolus value before entering bolus interval or # of boluses/hr.

LCD Display: ALERT NO KEYPAD ACTIVITY

Alert: No Keypad Activity

Cause: The pump has been displaying a non-running screen for two minutes without any keys being pressed. The alert will recur every 2 minutes as long as there are no key presses.

Resolution: Complete the activity for the displayed menu. The pump may also be turned off, however, all programming that was not complete will be discarded.

If the pump was running and the Options menu displayed when the alert occurred, the run screen for the therapy will be displayed when the alert is cleared. Any changes to the Options menu are discarded and will have to be re-entered.

LCD Display: ALERT NO THERAPY ENABLED

Alert: No Therapy Enabled

Cause: Attempting to accept the Enable Therapy menu with all therapies set to "OFF". At least one therapy must be enabled or "ON" for menu to be accepted.

Resolution: Enable the therapy or therapies that are to be active for the pump.

LCD Display: ALERT NOT ALLOWED, DOSE IN PROGRESS

Alert: Change is not allowed while a dose is in progress

Cause: Modification to the current field is not allowed while the dose is in progress.

Resolution: Wait until the dose in progress completes before modifying field.

LCD Display: ALERT NOT in PCA. NO BOLUS ALLOWED

Alert: Bolus requested when therapy is not PCA

Cause: Remote Bolus button or BOLUS key is pressed when not in PCA therapy.

Resolution: Remove the remote bolus cord from the pump if connected. Refrain from pressing the BOLUS key.

LCD Display: ALERT NUMBER OF DOSES INVALID

Alert: Number of doses entered exceeds maximum for therapy

Cause: The number of doses per bag in Intermittent therapy exceeds the allowable maximum.

Resolution: Recheck data entry. The number of doses requested must be less than or equal to the maximum for the therapy.

LCD Display: ALERT PERFORM ROUTINE MAINTENANCE

Alert: Maintenance Date (displayed only if date has arrived when pump is turned on)

Cause: Current Date is beyond the Maintenance Date.

Resolution: Return pump for routine maintenance.

LCD Display: ALERT RAMP INVALID FOR AMT/RATE

Alert: Ramp time entered is invalid for amt/rate already programmed

Cause: Ramp value entered for TPN is unacceptable given the desired rate and amount programmed.

Resolution: Recheck all data entered for prescription. If programming a new TPN therapy ("NEW PROGRAM" selected) and the ramp time is a requirement, enter the ramp time(s) and total time first followed by the next governing parameter (rate or volume to be infused).

LCD Display: ALERT RATE EXCEEDS DELTA RATE/DELTA TIME

Alert: Rate entered exceeds the delta time limits

Cause: The rate entered falls outside of the allowable range for the delta rate / delta time limits programmed.

Resolution: Recheck data entry or wait more time elapses then attempt to change the rate. Rate selected cannot exceed the range placed on medLIMITS.

LCD Display: ALERT RUNNING, PAUSE PUMP BEFORE OFF

Alert: Pause Infusion before turning off

Cause: The OFF key was pressed while the pump was infusing or a running screen displayed.

Resolution: Press "PAUSE" key before pressing "OFF" key to stop an infusion.

LCD Display: ALERT Rx IN PROGRESS, YES to Repeat – NO to Return

Alert: REPEAT Rx selected while a therapy is already in progress

Cause: REPEAT Rx was selected and there is a therapy currently in progress.

Resolution: To terminate the current therapy in progress, press YES. Resume will no longer be available. The Select Therapy menu will be displayed (if only one therapy is in programmed, the pre Rx menu for the therapy will be displayed). When the therapy to repeat is chosen, the last prescription data programmed will appear in the menu, excluding Delayed Start and Load Dose (if therapy was previously started). When the infusion is started the entire therapy will be repeated.

To continue the current therapy in progress, press NO. The Pause menu will be displayed.

LCD Display: ALERT Rx IN PROGRESS, YES New Program – NO to Return

Alert: NEW PROGRAM was selected while a therapy is already in progress

Cause: NEW PROGRAM was selected and there is a therapy currently in progress.

Resolution: To terminate the current therapy in progress, press YES. Resume will no longer be available. The Select Therapy menu will be displayed (if only one therapy is in programmed, an alert screen informing that the Rx will be erased will be displayed). When the therapy for the new program is chosen, entry of a new prescription will be required.

To continue the current therapy in progress, press NO. The Pause menu will be displayed.

LCD Display: ALERT Rx Programmed, YES to Erase – NO to Return

Alert: NEW PROGRAM was selected for a therapy that has an existing prescription

Cause: NEW PROGRAM was selected for a therapy that has an existing prescription.

Resolution: To erase the prescription for the selected therapy, press YES. The pre Rx menu will be displayed and entry of a new prescription will be required.

To preserve the prescription for the selected therapy, press NO. The Pause menu will be displayed.

LCD Display: ALERT SET NOT INSTALLED

Alert: Set Not Installed

Cause: Administration set is not properly installed.

Resolution: Install the appropriate **Curlin Infusion** administration set for the desired therapy. Use *ONLY* **Curlin Infusion** administration sets. See Chapter 1, figures 1.9 and 1.10 of the User's Manual for proper installation of the administration set.

LCD Display: ALERT SYSTEM CLOCK NOT SET

Alert: System Clock Not Set

Cause: The clock for the pump is not set

Resolution: Use the BIOMED SETUP menu Date and Time option to set the date and time.

LCD Display: ALERT THERAPY TIME INVALID

Alert: Invalid therapy time calculated

Cause: While programming or modifying an Intermittent therapy, the combination of dose duration and dose frequency result in a prescription with a duration greater than 288:00 HH:MM.

Resolution: Recheck data entry. The changes to the field must comply with the total time limit.

LCD Display: ALERT TIME LESS THAN RAMP TIME

Alert: Unable to enter a TPN therapy time that is less than the ramp times

Cause: Time value entered for TPN infusion is less than ramp times.

Resolution: Recheck data entry. TPN infusion time cannot be less than specified ramp times.

LCD Display: ALERT USING 2.0 ML DEFAULT, PRESS YES TO OFF FOR OFF

Alert: The air sensitivity currently being used by the pump is 2.0ml

Cause: YES was pressed at the "ACCEPT?" field on the Options menu and the displayed setting in Options for the AIR SENS field is OFF, however, the pump is using a setting of 2.0ml. This occurs when YES was not pressed when the "AIR SENS OFF. Using In Line Filter?" alert screen was last displayed and the down scroll arrow was used instead of the YES key to pass the AIR SENS field on the Options menu. Alert is only displayed when Options is displayed and pump is running.

Resolution: The cursor will be on the AIR SENS field when the alert is cleared. Press YES to display the "AIR SENS OFF. Using In Line Filter?" screen. Press YES if an air eliminating filter is being used. Press NO if no air eliminating filter is being used.

LCD Display: ALERT VALUE PREVENTS A SUBSEQUENT DOSE

Alert: Value entered will prevent the ability to program any more doses

Cause: The value entered for the Variable Dose Amt or Time field will cause no remaining amount or time for a subsequent.

Resolution: Recheck data entry.

If this is a new program, reprogramming of the therapy will be required if the error is in a previously programmed dose.

If the therapy is in progress or this is a repeat program that has not started and the error is in a previously programmed dose, turn the pump off and then on. Choosing to resume or repeat the therapy will display the last prescription that was completely programmed.

Alarms are conditions that require correction by the user and may be a safety issue but are not system errors of the pump. All alarm conditions stop the infusion. The only exception to this rule is at the "Infusion Complete" alarm, which allows the KVO rate, when programmed, to continue infusing. When an alarm occurs, a message is displayed on the screen, the audio alarm is sounded, and the red LED light flashes. The message stays on the screen until the pause key is pressed. The "Silence" key can be pressed to disable the audio alarm for one-minute intervals. The cause of the alarm must be resolved, or the alarm will recur when the infusion is restarted. To exit an alarm screen, press the "PAUSE" key, resolve the alarm condition, and then resume the therapy by selecting "RESUME" from the Pause Menu.

The following is a selected and alphabetical list of the alarms, conditions that cause the alarms, messages displayed on the screen, and actions necessary to resolve them:

LCD Display: ALARM AIR IN LINE
Alarm: Air In Line
Cause: Pump has detected air in the administration set during an infusion.
Resolution: Check air settings and remove air from administration set and IV bag or reservoir per protocol. (Disconnect from patient's access site before using prime function.)

LCD Display: ALARM DOOR OPEN or SET NOT PROPERLY INSTALLED
Alarm: Door is opened or administration set is not properly installed while the pump is running
Cause: Door is opened or administration set is not properly installed while the pump is running
Resolution: Check the door latch assembly and securely close the door. If the door was closed properly, re-install the set according to the installation instructions. See Figure 1.8 for instructions on closing door.

LCD Display: ALARM DOWN OCCLUSION
Alarm: Occlusion between pump and patient access site
Cause: Pump has detected excessive pressure in the administration set between the pump and the patient's access site.
Resolution: Check administration set from pump to patient access site for potential causes of occlusion. If tubing is kinked, straighten it. If filter is blocked , replace administration set. If patient access is obstructed, check it according to protocol. Consider changing the down occlusion pressure setting from Low to High. If the occlusion is cleared without the need to open the pump door, the pump automatically resumes its operation. Note: If the door of the pump is opened to clear the occlusion, (e.g. replacing the set), you must press the pause key, select "Resume," and then press the "RUN" key to restart the infusion.

LCD Display: ALARM EMPTY BATTERY

Alarm: Empty Battery

Cause: The "C" cell battery no longer has sufficient power to run the pump. Alarm will not be displayed when external battery pack is depleted.

Resolution: Turn the pump off. Replace both of the "C size" batteries. Recharge or replace the battery pack if applicable.

LCD Display: ALARM HIGH UP PRESSURE

Alarm: High positive pressure between the pump and IV reservoir

Cause: The pump has detected excessive positive pressure between the pump and the IV Source.

Resolution: Press Pause to clear alarm screen. Check for excessive pressure on the IV bag and relieve it. When ready to begin infusion, select RESUME.

LCD Display: ALARM REPLACE SET 3

Alarm: Replace Set

Cause: The pump has detected that the disposable administration set must be replaced.

Resolution: Replace the disposable administration set, prime the set if necessary, and when ready to begin infusing, select RESUME to continue the infusion.

LCD Display: ALARM REPLACE SET 4

Alarm: Replace Set

Cause: The pump has detected excessive pressure in the disposable administration set.

Resolution: Make sure that all the IV clamps are open. Relieve pressure, open and close pump door.

LCD Display: ALARM SET NOT INSTALLED

Alarm: Administration Set Not Installed

Cause: The administration set has not been installed or has been installed improperly.

Resolution: Install appropriate **Curlin Infusion** Administration set for desired therapy. Use ONLY **Curlin Infusion** Administration sets. If the blue tubing guide on the administration set has been damaged in any way, replace the entire administration set. See Chapter entitled *Introduction*, Figures 1.9 and 1.10 for proper installation of administration set.

LCD Display: ALARM UNATTENDED PUMP

Alarm: No key presses have occurred while the pump was paused

Cause: When a therapy is in progress and a pump is placed in a paused state and left unattended for more than two minutes, this alarm will sound. If the prime menus are left unattended for one minute, this alarm will sound

Resolution: Press the "PAUSE" key to clear the alarm screen. If required, return to the screen displayed at the time of the alarm and complete the activity that was in progress, or resume the infusion, or turn the pump off.

LCD Display: ALARM UP OCCLUSION

Alarm: Negative pressure detected between the pump and IV reservoir

Cause: Pump has detected excessive negative pressure in the administration set between the medication reservoir or IV bag ("up stream") and the pump.

Resolution: Pause the pump. Check administration set from IV bag to pump for potential causes of occlusion. Open the door of the pump and remove the administration set from the pumping chamber. If an area of the tubing is flattened, massage that area to a more rounded shape. If a clamp is in place, open it; if tubing is kinked, straighten it. Replace administration set back in the pump. When ready to begin infusing, select RESUME. If the alarm continues, use a new administration set or if the IV bag is almost empty, replace it.

LCD Display: CMS NOT ENABLED

Alarm: Pump is not CMS capable

Cause: A CMS system had attempted to communicate with this pump which was not CMS capable

Resolution: This pump is not capable of CMS communication and should not be used for any CMS function.

LCD Display: COMM TIMEOUT

Alarm: Communication Timeout occurred during Pump/CMS communication

Cause: The communication link between the pump and the P.C. running CMS was broken (i.e., the cable between them was unplugged) during a CMS upload to the pump

Resolution: Check the cable connection between the pump and the PDA or P.C.

LCD Display: INFUSION COMPLETE

Alarm: Infusion Complete

Cause: Programmed "Amount to be Infused" is depleted to 0.

Resolution: If the therapy is to be repeated, press "PAUSE", add another IV bag or reservoir and select the "REPEAT Rx" function.

If the therapy will not be repeated, press "PAUSE" key and turn the pump off by pressing the "OFF" key.

LCD Display: UPLOAD CHAR ERR

Alarm: While uploading a prescription to the pump a character was misinterpreted

Cause: Faulty communications link between the pump and the P.C. running CMS

Resolution: Replace the communication cable between the pump and the PDA or PC. If other programs are running on the PC that might with the operation of CMS/PC, close those programs.

LCD Display: UPLOAD CkSum ERR

Alarm: CMS Upload experienced a Check Sum Error

Cause: At the completion of an upload of a CMS script the Check Sum signature from the CMS system did not match the signature calculated by the pump

Resolution: A second attempt to upload the script should be made. If the second attempt fails, then a new script should be created for the desired prescription. (See CMS User's Manual).

ERROR CODES

If a pump goes into an error code, a numeric code will appear on the display screen. If this has occurred, turn the pump off and back on. If the error code message does not reappear, continue using the pump.

If the pump passes its start-up diagnostic tests, resume the therapy. If an error code recurs, turn the pump off, replace it, and report the error code to the healthcare provider**.**

Error Code	LCD Displayed Message	Probable Cause	Corrective Action
10	Turn pump OFF/ON or Replace Battery or Call Provider	Motor Control Loss	▪ Replace batteries. ▪ Connect AC Adaptor. ▪ Turn power off/on and resume.
11	Turn pump Off/On	Keyboard Buffer Full	Report error code to factory.
12	Air Sensor Turn pump Off/ON or Call provider	Air In Line test circuit failure	Report error code to factory.
13	Pressure Sens DN Check Admin Set turn pump Off/On	Bad Down Strain Beam	Replace set and Resume.
14	Pressure Sens UP Check Admin Set Turn pump Off/On	Bad Up Strain Beam	Replace set and Resume.
15	Turn pump Off/On or Call Provider	Bad Encoder	Report error code to factory.
16	Turn pump Off/On or call Provider	Bad Analog to Digital Conversion	Report error code to factory.
17	Turn pump OFF Call Provider	Bad Clock	Report error code to factory.
18	Call Provider	Bad Air In Line comparator register reading from FPGA.	Report error code to factory.
19	Turn Pump OFF Call Provider	Bad Door Sensor	Report error code to factory.
20	Empty Lithium Battery Call Provider	Empty Lithium Battery	Report error code to factory.
21	Turn pump Off Call Provider	Bad Tristate Pin	Report error code to factory.
22	Turn Pump OFF Call Provider	Bad Test Pin	Report error code to factory.
23	Turn Pump OFF Call Provider	Bad Reverence Voltage	Report error code to factory.
24	Turn Pump OFF Call Provider	Bad Motor Supply	Report error code to factory.
25	Stalled Pump Turn pump OFF/ON Call Provider	Broken Motor Shaft	Turn power off/on and resume.
26	Turn Pump OFF Call Provider	Bad Motor Inhibit	Report error code to factory.
27	Turn Pump OFF Call Provider	Bad Motor Supply Enable	Report error code to factory.
28	Turn Pump OFF Call Provider	Bad Motor Logic Enable	Report error code to factory.
29	Low Alarm Batt Connect AC Adapt Turn pump OFF/ON	Low Auxiliary Battery	Connect pump to AC Battery Eliminator and turn power off/on to pump. May require turning pump off/on a few times.
30	Turn Pump OFF Call Provider	Bad Auxiliary Battery	Report error code to factory.
31	Turn Pump OFF Call Provider	Audio Main Speaker Low	Report error code to factory.
32	Turn Pump OFF Call Provider	Audio Main Speaker High	Check/replace D31. Replace PCB.
33	Low Alarm Audio Connect AC Adapt Turn pump OFF/ON	Audio Auxiliary Speaker Low	Connect pump to AC Battery Eliminator and turn power off/on to pump. May require turning pump off/on a few times.

Error Code	LCD Displayed Message	Probable Cause	Corrective Action
34	Turn Pump OFF Call Provider	Audio Auxiliary Speaker High	Report error code to factory.
35	History File Turn pump OFF/ON	History Log Erased	Report error code to factory.
36	Bad Cal Data Turn pump OFF Call Provider	Bad EEPROM Data CRC value	Report error code to factory.
37	Turn Pump OFF Call Provider	Analog to Digital Converter did not Run	Report error code to factory.
38	Turn Pump OFF Call Provider	Air In Line Circuit not Tested	Report error code to factory.
39	Turn Pump OFF Call Provider	Bad EEPROM Data CRC 2	Report error code to factory.
40	Turn Pump OFF Call Provider	Bad EEPROM Data Header	Report error code to factory.
41	Bad Cal Range Turn Pump OFF Call Provider	Bad EEPROM Limit	Report error code to factory.
45	Turn pumpOFF/ON Call Provider	Motor Control Loss	Report error code to factory.
46	Turn pumpOFF/ON Call Provider	Motor Control Loss	Report error code to factory.
100	Turn pumpOFF/ON Call Provider	Menu not found	Report error code to factory.
101	Turn pumpOFF/ON Call Provider	Mode not found	Report error code to factory.
102	Turn pumpOFF/ON Call Provider	Item not found	Report error code to factory.
103	Turn pumpOFF/ON Call Provider	Field not found	Report error code to factory.
104	Turn pumpOFF/ON Call Provider	Data mismatch	Report error code to factory.
105	Turn pumpOFF/ON Call Provider	No menu pointer	Report error code to factory.
106	Turn pumpOFF/ON Call Provider	Count out of range	Report error code to factory.
107	Turn pumpOFF/ON Call Provider	Display task queue overflow	Report error code to factory.
108	Turn pumpOFF/ON Call Provider	Divide by zero	Report error code to factory.
109	Turn pumpOFF/ON Call Provider	Concentration zero value	Report error code to factory.
110	Turn pumpOFF/ON Call Provider	Invalid occurrence	Report error code to factory.
111	Turn pumpOFF/ON Call Provider	Multiple param struct access	Report error code to factory.
112	Turn pumpOFF/ON Call Provider	Multiple status struct access	Report error code to factory.
140	Turn pumpOFF/ON Call Provider	Infusion volume out of range	Report error code to factory.
141	Turn pumpOFF/ON Call Provider	Infusion rate out of range	Report error code to factory.
142	Turn pumpOFF/ON Call Provider	Infusion time out of range	Report error code to factory.
143	Turn pumpOFF/ON Call Provider	Volume conversion error	Report error code to factory.
144	Turn pumpOFF/ON Call Provider	Rate conversion error	Report error code to factory.
145	Turn pumpOFF/ON Call Provider	Time conversion error	Report error code to factory.
146	Turn pumpOFF/ON Call Provider	Bad PCA dose ratio	Report error code to factory.
160	Stuck PERIOD Key Turn pump OFF/ON or Call Provider	Stuck Period / Silence Key	If key was pressed, release it. Otherwise report error code to factory.
161	Stuck ON OFF Key Turn pump OFF/ON or Call Provider	Stuck On / Off Key	If key was pressed, release it. Otherwise report error code to factory.

Error Code	LCD Displayed Message	Probable Cause	Corrective Action
162	Stuck 0 Key Turn pump OFF/ON or Call Provider	Stuck 0 Key	If key was pressed, release it. Otherwise report error code to factory.
163	Stuck 1 Key Turn pump OFF/ON or Call Provider	Stuck 1 Key	If key was pressed, release it. Otherwise report error code to factory.
164	Stuck 2 Key Turn pump OFF/ON or Call Provider	Stuck 2 Key	If key was pressed, release it. Otherwise report error code to factory.
165	Stuck 3 Key Turn pump OFF/ON or Call Provider	Stuck 3 Key	If key was pressed, release it. Otherwise report error code to factory.
166	Stuck 4 Key Turn pump OFF/ON or Call Provider	Stuck 4 Key	If key was pressed, release it. Otherwise report error code to factory.
167	Stuck 5 Key Turn pump OFF/ON or Call Provider	Stuck 5 Key	If key was pressed, release it. Otherwise report error code to factory.
168	Stuck 6 Key Turn pump OFF/ON or Call Provider	Stuck 6 Key	If key was pressed, release it. Otherwise report error code to factory.
169	Stuck 7 Key Turn pump OFF/ON or Call Provider	Stuck 7 Key	If key was pressed, release it. Otherwise report error code to factory.
170	Stuck 8 Key Turn pump OFF/ON or Call Provider	Stuck 8 Key	If key was pressed, release it. Otherwise report error code to factory.
171	Stuck 9 Key Turn pump OFF/ON or Call Provider	Stuck 9 Key	If key was pressed, release it. Otherwise report error code to factory.
172	Stuck UP Key Turn pump OFF/ON or Call Provider	Stuck Up Arrow Key	If key was pressed, release it. Otherwise report error code to factory.
173	Stuck NO Key Turn pump OFF/ON or Call Provider	Stuck No / Change Key	If key was pressed, release it. Otherwise report error code to factory.
174	Stuck YES Key Turn pump OFF/ON or Call Provider	Stuck Yes / Enter Key	If key was pressed, release it. Otherwise report error code to factory.
175	Stuck DOWN Key Turn pump OFF/ON or Call Provider	Stuck Down Arrow Key	If key was pressed, release it. Otherwise report error code to factory.
176	Stuck RUN Key Turn pump OFF/ON or Call Provider	Stuck Run / Pause Key	If key was pressed, release it. Otherwise report error code to factory.
177	Stuck HELP Key Turn pump OFF/ON or Call Provider	Stuck Help / Options Key	If key was pressed, release it. Otherwise report error code to factory.
178	Stuck PRIME Key Turn pump OFF/ON or Call Provider	Stuck Prime / Bolus Key	If key was pressed, release it. Otherwise report error code to factory.
179	Bolus Cord Unplug Cord Turn pump OFF/ON	Stuck Remote Bolus Switch	If Bolus Switch was pressed, release it. Replace Bolus assembly. Otherwise report error code to factory.
180	Turn pump OFF/ON Call Provider	Invalid Key	Report error code to factory.
200	Turn pump OFF/ON Call Provider	Invalid system state	Report error code to factory.
201	Turn pump OFF/ON Call Provider	Bad power up test number	Report error code to factory.
202	Turn pump OFF/ON Call Provider	Bad power up state	Report error code to factory.
203	Turn pump OFF/ON Call Provider	Bad A to D Max address	Report error code to factory.

Error Code	LCD Displayed Message	Probable Cause	Corrective Action
Error Code	LCD Displayed Message	Probable Cause	Corrective Action
204	Turn pump OFF/ON Call Provider	Bad run state	Report error code to factory.
205	Turn pump OFF/ON Call Provider	Bad pump status	Report error code to factory.
206	Turn pump OFF/ON Call Provider	Bad volume type	Report error code to factory.
207	Turn pump OFF/ON Call Provider	Bad rate type	Report error code to factory.
208	Turn pump OFF/ON Call Provider	Bad motor test number	Report error code to factory.
209	Turn pump OFF/ON Call Provider	Bad hist type	Report error code to factory.
210	Turn pump OFF/ON Call Provider	Bad EEPROM Data ID	Report error code to factory.
211	Turn pump OFF/ON Call Provider	Bad run time CRC state	Report error code to factory.
212	Turn pump OFF Call Provider	Bad EEPROM Write	Report error code to factory.
213	Turn pump OFF Call Provider	Bad total volume	Report error code to factory.
214	Turn pump OFF Call Provider	Bad total time	Report error code to factory.
300	Turn pump OFF Call Provider	Stack Overflow	Report error code to factory.
301	Turn pump OFF Call Provider	Bad Flash	Report error code to factory.
302	Turn pump OFF Call Provider	Bad Run Time Flash	Report error code to factory.
303	Turn pump OFF Call Provider	Bad Ram	Report error code to factory.
304	Turn pump OFF Call Provider	Bad output data	Report error code to factory.
305	Turn pump OFF Call Provider	Bad active data	Report error code to factory.
306	Turn pump OFF Call Provider	Bad overflow count	Report error code to factory.
307	Turn pump OFF Call Provider	Therapy data CRC failure	Report error code to factory.
308	Watch Dog Call Provider	Bad Watch Dog Test	Report error code to factory.
309	Turn pump OFF Call Provider	Stopped Motor Motion Count	Report error code to factory.
310	Turn pump OFF Call Provider	Stopped Motor Motion Volt	Report error code to factory.
311	Turn pump OFF Call Provider	Bad Active Flags	Report error code to factory.
312	Turn pump OFF Call Provider	Bad Active Flags 2	Report error code to factory.
313	Turn pump OFF Call Provider	Negative Motor Motion	Report error code to factory.
314	Turn pump OFF Call Provider	Therapy params write violate	Report error code to factory.
315	Turn pump OFF Call Provider	Therapy status write violate	Report error code to factory.
316	Turn pump OFF/ON Call Provider	Configuration settings may have been compromised	Try to turn off pump then on. If OK, continue, if error recurs, return for service.
317	Turn pump OFF/ON Call Provider	Configuration settings may have been compromised	Try to turn off pump then on. If OK, continue, if error recurs, return for service.

Chapter

8

Accessories

There are a number of helpful accessories for use with the ***Curlin Infusion*** *pumps. Some of the most common accessories are described in this chapter. For a complete and updated list, refer to the Moog website. www.moog.com/medical.*

Detachable Pole Clamps

The pole clamps attach to the Lockable Safety Shells or the Pump Holster and can accommodate a pole of .75 to 1.25-inch diameter. There is a **Standard Pole Clamp (340-0104)** as well as a **Lockable Pole Clamp (340-0125)** available.

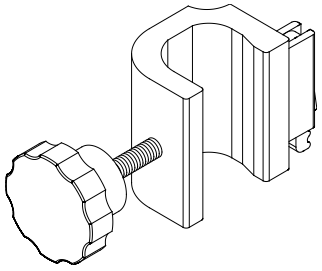


Figure 8.1 Detachable Pole Clamp

Remote Bolus Cord (340-2005)

This five-foot flexible cable offers a remote push button switch for the patient's use in administering bolus dosing in the PCA therapy. This accessory cannot be used in any therapy except PCA; and if it is left connected for any other therapy, an alert message will display on the pump screen, instructing the operator to remove this cable.

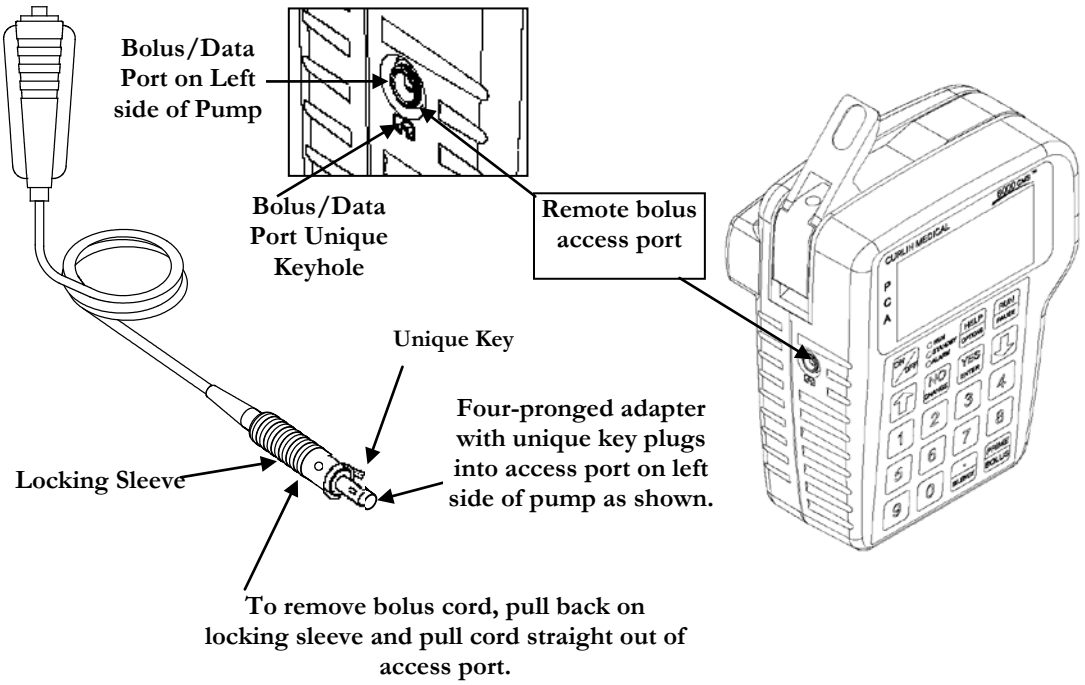


Figure 8.2 Remote Bolus Cord

Lockable Safety Shells

These accessories are hard-plastic, durable cases that are **lockable with a key**. The **250 ml safety shell (360-0018)** will secure medication bags of up to 250 ml, administration set and the pump, and yet allow access to the pump's keypad, controls, communication connectors, and battery compartment door.

The 250 ml **safety shells** will also secure syringes and is available in light blue.

All security shells can be pole mounted with the detachable pole clamp, set on a stable flat surface, or carried in a soft carry pack.

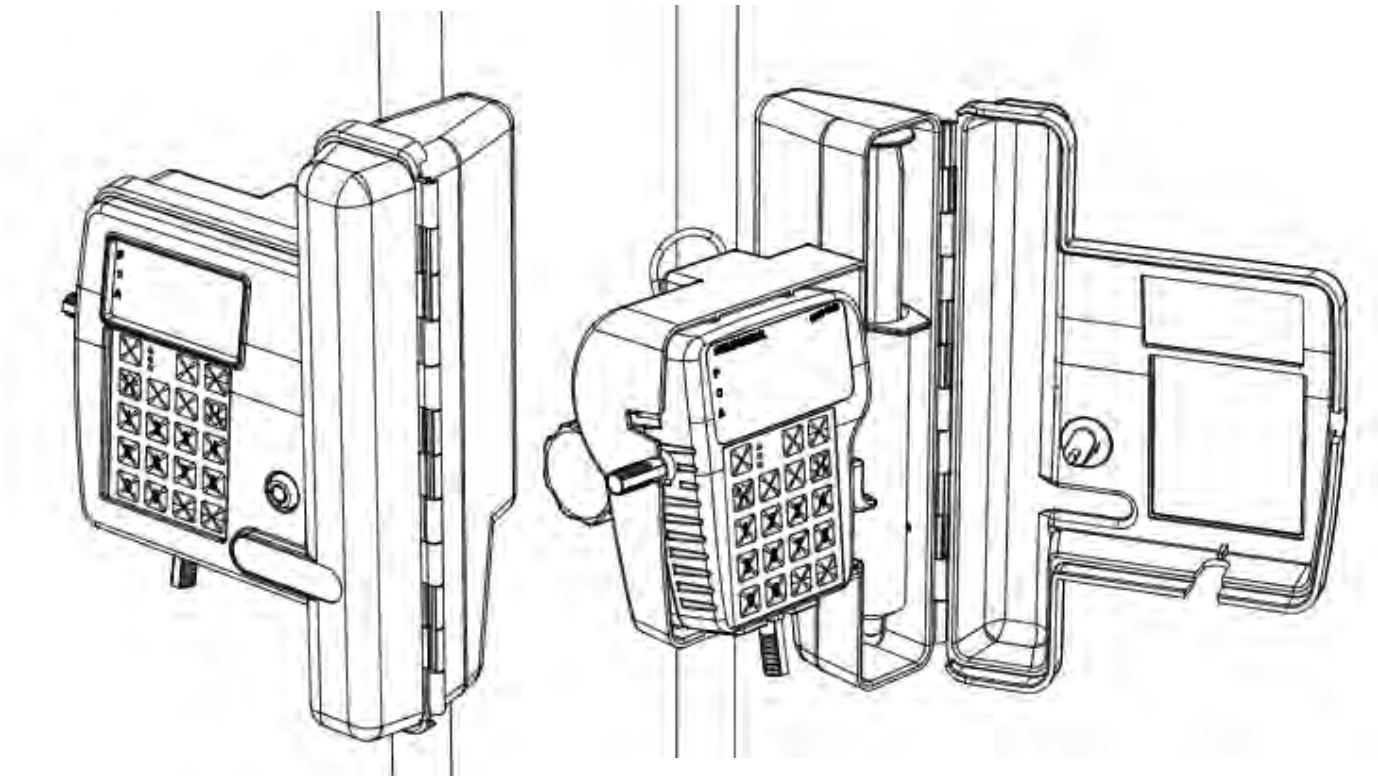


Figure 8.3 Safety Shells

Additional graphics below show how to install the Pole Clamp on the Lock Box.

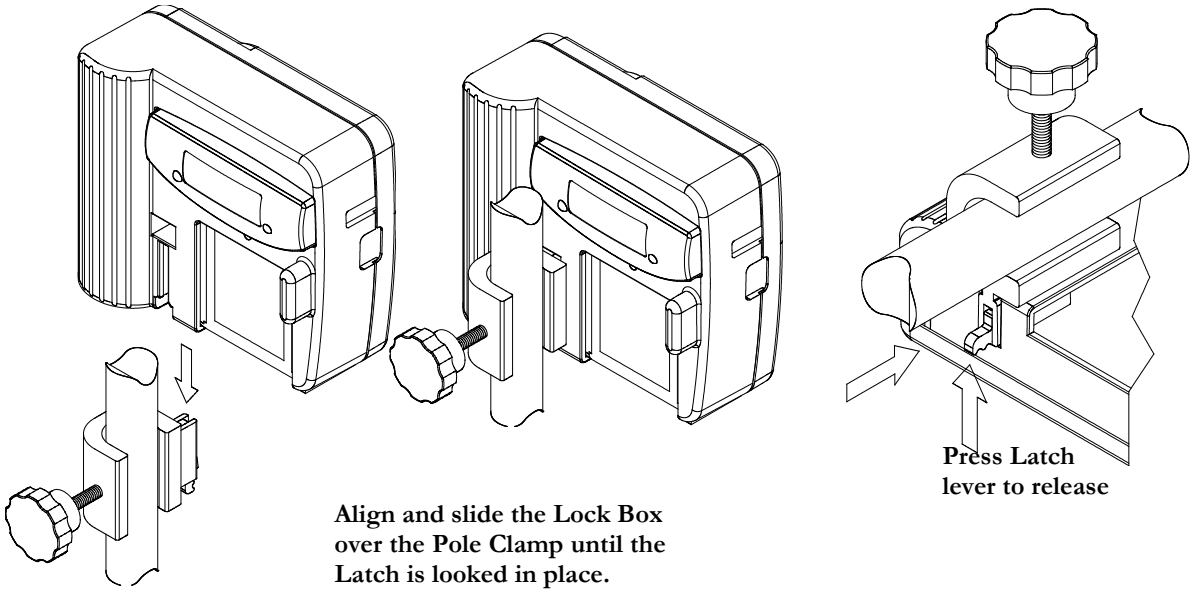
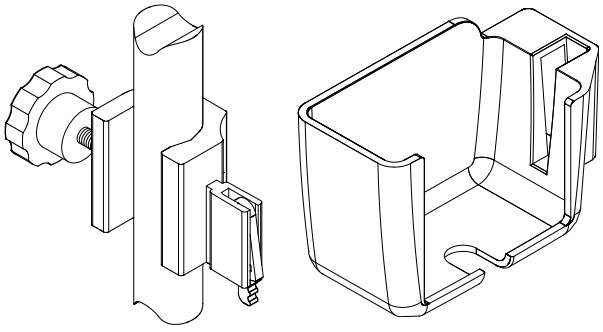


Figure 8.4 Pole Clamp and Safety Shell Installation

Pump Holster (340-0111)

This accessory is a simple "slide pump in" case with a pole clamp-mounting bracket on the back, allowing the pump to securely be mounted onto a pole and yet allow the user easy access to the display, keypad, controls, battery compartment, and connectors for external power or communications.



Curlin Medical Holster with pole mount to be attached to an IV pole. The back of the holster slides onto the pole mount and locks securely in place.

Figure 8.5 Pump "Holster"

Keypad Covers

These accessories are hard-plastic, durable keypad covers that can optionally be used with the (340-0141, 340-0141-01 and 340-0141-02) lockboxes to cover the keypad. The keypad covers are available in four configurations:

- **Type 0 (340-0142)** covers all the keys;
- **Type I (340-0142-01)** covers all the keys, with the exception of the "SILENCE" and "RUN/PAUSE" keys;
- **Type II (340-0142-02)** covers only the numerical and "PRIME/BOLUS" keys.
- **Type III (340-0142-03)** covers all the keys, with the exception of the "SILENCE" key.

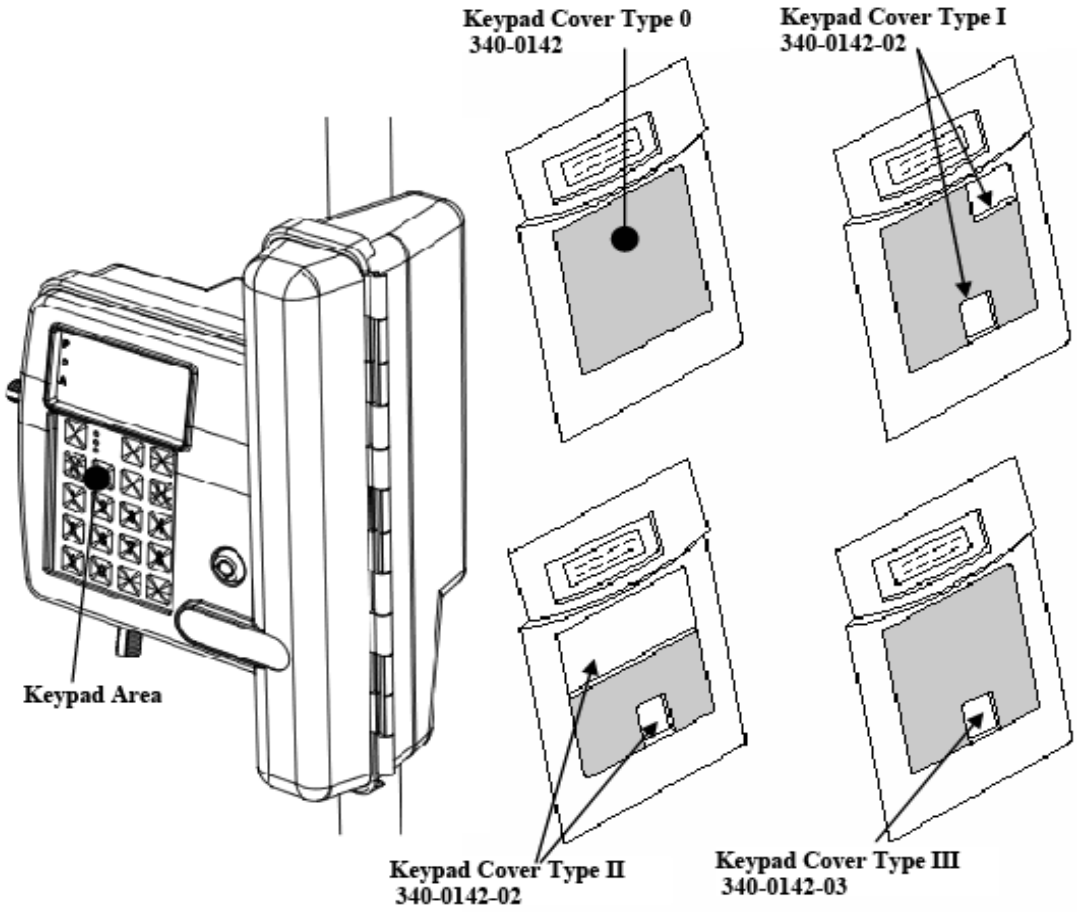


Figure 8.6 Keypad Covers

Cleaning, Environmental Conditions, and Maintenance

Following the guidelines for care listed below will assure the best appearance and longest life of the ***Curlin Infusion*** pump.

Cleaning

The **Curlin Infusion** pumps are designed to be resistant to fluid spillage from any direction but are *not* to be immersed in solutions. Keep the pump and the pumping surfaces clean, dry, and free of fluid spillage at all times. Do not use the pump in the shower, sauna, or steam bath and do not position the pump where it could accidentally be dropped into a container of fluid, (e.g. basin, tub, or toilet.) The pumping surfaces can be cleaned with a cloth dampened with any of the following solutions, and dried with a soft cloth without harming the pump.

Clean the pump, remote bolus cord, AC adapter, safety shells, holster and battery pack with a soft cloth dampened in any of the following:

- Warm soapy water (do not submerse)
- Isopropyl alcohol
- Household bleach, diluted 9:1 with water
- Commercial disinfectant

(Do NOT spray the cleaning solution directly on the pump.)

IMPORTANT INFORMATION

The following cleaning/disinfecting methods are specifically prohibited:

- **Do not use full strength bleach**
- **Do not autoclave or use steam sterilization**
- **Do not clean in a dishwasher**
- **Do not use acetone, lacquer thinner, or any other solvents or abrasive cleansers to clean the pump**

With proper care, the pump will retain its finish for many years of service.

Transport and Storage

During transport or storage, the batteries should be removed; the pump should be cleaned and kept in its hard-shell packing case. The environmental conditions should be within the range of:

- An ambient temperature -20° F to 140°F
- A relative humidity range of 10% to 95%
- An atmospheric pressure range of 500 Hpa to 1060 Hpa

All accessories should be stored in their original packing containers to ensure the longest service.

Operating Environment

The following conditions of ambient temperature range and relative humidity are recommended for the fluid delivered accuracy of:

- Temperature: ±5% Accuracy: 55 to 90°F
  ±10% Accuracy: 45 to 55°F
  ±10% Accuracy: 90 to 104°F
- Relative Humidity: 15 to 95% Non-condensing.

Maintenance

Yearly preventive maintenance checks are recommended and will be prompted in the "Set-Up" menu and on the display when the date occurs. This date accrues from the date of factory release and is reset when each annual maintenance check is completed.

Any time an error code message occurs repeatedly or any time a pump appears to be tampered with or shows any indication of not performing to standards, the pump should be returned to the healthcare provider dispensing the pump.

At no time should unauthorized personnel open the hard case of the pump.

Limited Warranty

ZEVEX, Inc. (the "Manufacturer") warrants to the Original Purchaser that the infusion pump (the "Pump"), not including accessories, shall be free from defects in material and workmanship under normal use, if used in accordance with this User's Manual, for a period of one year from the actual date of sale to the Original Purchaser. THERE ARE NO OTHER WARRANTIES.

The warranty does not cover normal wear and tear and maintenance items, and specifically excludes batteries, administration sets, extension sets or any other accessory items or equipment used with the Pump.

Subject to the conditions of and upon compliance with this Limited Warranty, the Manufacturer will repair or replace at its option without charge (except for a minimal charge for postage and handling) any Pump (not including accessories) which is defective if a claim is made during such one year period.

The following conditions, procedures, and limitations apply to the Manufacturer's obligation under this warranty:

1 PARTIES COVERED BY THIS WARRANTY: This warranty extends only to the Original Purchaser of the Pump. This warranty does not extend to subsequent purchasers. The Original Purchaser may be a patient, medical personnel, or a medical facility that purchases the Pump for treatment of patients. The Original Purchaser should retain the invoice or sales receipt as proof as to the actual date of purchase.

2 WARRANTY PERFORMANCE PROCEDURE: Notice of the claimed defect must be made in writing or by telephone to the Manufacturer as follows: **Customer Service Department, ZEVEX, Inc., 4314 ZEVEX Park Lane, Salt Lake City, UT 84123, (801) 264-1001**. Notice to the Manufacturer must include date of purchase, model and serial number, and a description of the claimed defect in sufficient detail to allow the Manufacturer to determine and facilitate any repairs that may be necessary. AUTHORIZATION MUST BE OBTAINED PRIOR TO RETURNING THE PUMP, and confirmed with issuance of an RMA (Return Materials Authorization) number. When authorized, the Pump must be properly and carefully packaged and returned to the Manufacturer, postage prepaid. Any loss or damage during shipment is at the risk of the sender.

3 CONDITIONS OF WARRANTY: The warranty is void if the pump has been 1) repaired by someone other than the Manufacturer or its authorized agent; 2) altered so that its stability or reliability is affected; 3) misused; or, 4) damaged by negligence or accident. Misuse includes, but not limited to, use not in compliance with the User's Manual or use with non approved accessories. The pump is a sealed unit, and the fact that the seal has been broken will be considered conclusive evidence that the Pump has been altered or misused. Removal or damage to the Pump's serial number will invalidate this warranty.

4 LIMITATIONS AND EXCLUSIONS: Repair or replacement of the pump or any component part thereof is the EXCLUSIVE remedy offered by the Manufacturer. The following and limitations shall apply:

   a. No agent, representative, or employee of the Manufacturer has authority to bind the Manufacturer to any representation or warranty, expressed or implied.
   b. THERE IS NO WARRANTY OF MERCHANTABILITY OR FITNESS OR USE OF THE PUMP FOR ANY PARTICULAR PURPOSE.
   c. The pump can only be used under the supervision of medical personnel whose skill and judgment determine the suitability of the Pump for any particular medical treatment.
   d. All recommendations, information, and descriptive literature supplied by the Manufacturer or its agents are believed to be accurate and reliable, but do not constitute warranties.

The Manufacturer disclaims responsibility for the suitability of the pump for any particular medical treatment or for any medical complications resulting from the use of the Pump. The Manufacturer shall not be responsible for any incidental damages or consequential damages to property, loss of profits, or loss of use caused by any defect or malfunction of the pump.

This warranty gives the Original Purchaser specific legal rights, and the Original Purchaser may have other legal rights, which may vary from state to state.

Procedure for Checking Pumps Between Patients

Purpose

This procedure is provided for use by caregivers to ensure pumps meet all critical performance specifications prior to use of the product on new patients. This procedure is for use by technicians to inspect, test, and prepare a pump for use by a new patient. The test checks for volumetric accuracy and all pump safety systems. Also, various parameters are initialized.

Note: Between patient testing is not required for Curlin Infusion pumps when anti-siphon valve administration sets are being used.

This procedure is intended to be used by technicians familiar with operation of the Curlin *PainSmart*™ IOD infusion pump.

Note: This procedure does not replace the recommended Annual Preventive Maintenance. The pump should receive preventive maintenance by a qualified technician at least annually.

Outpatient/Home Care/ Alternate Care Site Environment

If the pump is used in an outpatient, Home Care, or Alternate Care setting, the pump must be checked between patients using the following procedure.

Inpatient/Hospital Environment

Moog highly recommends that the hospital conduct in between patient checks using the following procedure. Should the hospital determine, for whatever reason, that the in between patient check is not practical, Curlin Medical recommends performing volumetric accuracy checks at regular intervals, to ensure the pump continues to meet specifications.

Required Equipment

The following equipment is required for this procedure:

- A NEW Administration Set (old or used sets are not suitable for the volumetric delivery and pressure measurements tests.)
- Volumetric measurement device (i.e. calibrated weight scale, burette or syringe.)
- Beaker with clean, fresh water or I.V. container.

Note: If the administration set contains a filter, sterile saline or sterile water should be used.

Maintenance Between Patients

1 CLEAN AND INSPECT PUMP FOR DAMAGE:

a. Clean the pump with a soft cloth dampened with a designated antiseptic, disinfectant solution or a solution of household bleach diluted 9:1 with water. Be sure to clean all surfaces, including the pumping chamber.

b. Inspect pump for damage. Check all case surfaces for cracks. Open the pumping chamber door. Examine the finger seal to confirm that no cuts or tears exist. Confirm that the door is well aligned by closing it with no unreasonable interference. Verify that the Power and Bolus/Data connectors are securely mounted on the case (no loose nuts). Verify that all labels are legible and in good condition. Check that Keypad is securely attached to case and that no corners are pealing up. Verify that

the Battery Door is in place and that the Battery Contacts in the Battery Compartment are not deformed.

Refer to the picture below when executing the following instructions.

2 Volumetric Delivery Test:

a. Install a new primed Administration Set in the pump and latch the door.

b. Connect the up-stream side of administration set to a water source 12 inches +/-2 inches above the pump.

c. Connect the downstream side of the administration set to a volumetric accuracy measurement device (i.e. a calibrated scale or burette) at the same level as the pump +/- 2 inches.

d. Program one of the following Infusions based on the default in the pump:

Pre Rx:		Pre Rx:	
UNITS:	ml	UNITS:	mg
Admin Rt:	EPI	CONC:	1 mg/ml
Load Dose:	0.0 ml	Admin Rt:	IV
MedLIMITS:	OFF	Load Dose:	OFF
NEXT?	YES	MedLIMITS:	OFF
		NEXT?	YES
Prescription:		Prescription:	
BAG VOL:	100 ml	BAG VOL:	100 ml
RATE:	0 ml/hr	RATE:	0 mg/ml
Pt Bolus:	9.9 ml	Pt Bolus:	9.9 mg
BOLS INT:	1 min	BOLS INT:	1 min
#BOLS/hr	1	#BOLS/hr	1
DONE?	YES	DONE?	YES

e. Start the pump by pressing RUN button.

f. Press the BOLUS key to start a bolus.

g. When the 9.9 ml infusion is over, observe the amount volume infused as measured on the measurement device. Passing measurement is: 9.4 g to 10.4 g (9.4 ml to 10.4 ml).

h. When Infusion complete occurs verify that the Audio Alarm beeps and that the Green LED is flashing.

3 CHECK ALL SAFETY SYSTEMS:

Repeat the same Rx as above. Start the infusion and press the BOLUS key to start a bolus dose.

a. **Low Down Stream Occlusion** - Go to the Option screen and confirm that the Down Occlusion level is set to LOW. Start the infusion and while the pump is running pinch off or close the down stream side of the administration set. Confirm the pump goes into Down Occlusion alarm. Open or clear the down stream occlusion and confirm the pump resumes infusion.

b. **High Down Stream Occlusion** - Go to the Option screen and change the Down Occlusion level to HIGH. Again pinch off or close the clamp on the down-stream side of the administration set. Confirm the pump goes into Down Occlusion. Open or clear the down stream occlusion and confirm the pump resumes infusion.

c. **Up Stream Occlusion** - Resume the infusion pumping then pinch off or close the up-stream side of the administration set. Confirm that the pump goes into an Up Occlusion alarm. Open or clear the up stream occlusion, press Pause then Resume the infusion.

d. **Door Sensor** - With infusion in progress open the door latch and observe ALARM DOOR OPEN OR SET NOT PROPERLY INSTALLED. Press Pause and then YES to RESUME on the Pause Menu.

e. **Set Sensor** - Remove the administration set from the pump and reinstall it so that the flow stop and tubing guide are not in the pumping chamber. Close the door latch and start an infusion by pressing the RUN key. Observe "ALERT DOOR OPEN

or SET NOT PROPERLY INSTALLED" will occur.

f. **Air In Line Sensor** - Reinstall the administration set properly and resume the infusion. From the Options menu set the AIL SENS (AIR IN LINE sensitivity) to 0.1ml. With the pump running introduce an Air Bolus of approximately 0.5ml into the fluid path at the input side of the pump. Once approximately 0.1ml of air has past the AIL detector, an ALARM AIR IN LINE will occur. Pause and Prime the air bolus past the AIL sensor, then resume the infusion.

g. **Auxiliary Alarm** - With the pump running observe the Vol INF (volume infused on the run screen) and disconnect the pump from all power sources (unplug the AC Adapter eliminator and remove one C-Cell battery.) Observe that the Auxiliary Alarm System is activated, with the Red LED and Beeper on constant (not intermittently beeping). With the battery still out of the pump, press the ON/OFF button to turn off the auxiliary alarm. Replace the battery in the battery compartment and turn on the pump. Resume the therapy and confirm that the infusion continues where it stopped due to Auxiliary Alarm occurrence.

4 SET BIOMED CLINICIAN MENU ITEMS:

a. Turn the pump Off and back On. While pump is performing Self Tests confirm that the pump is set to the correct Date and Time. If date and time are not correct, program the correct date and time from the Biomed/Clinician menu.

Note: Setting of the Date and Time is not allowed if 'RESUME' (infusion is still active) is available on the Pause menu. Use the CLEAR Pt Data/Rx option on the Biomed/Clinician menu to clear the prescriptions and end the infusion. The Date and Time can then be programmed.

b. From the Program/Biomed Setup Menu select Biomed Setup.

c. At the Enter Code Menu, enter the Clinician Code

d. (Optional) Erase the patient history file using the Clear Hx Log Menu.

e. Erase any previous Programmer ID, Patient ID, Rx Number, Invoice Number, Lock Level setting, and prescriptions using the Clear Pt Data/Rx Menu

f. Observe the Pump Maintenance Date, top of the Biomed/Clinician Menu, to insure adequate time for the next use by patient.

g. Sign, date and return the Check List to your record file. Pumps that pass can be used for new patients. Pumps that fail are to be returned for service.

Chapter

10

Technical Specifications

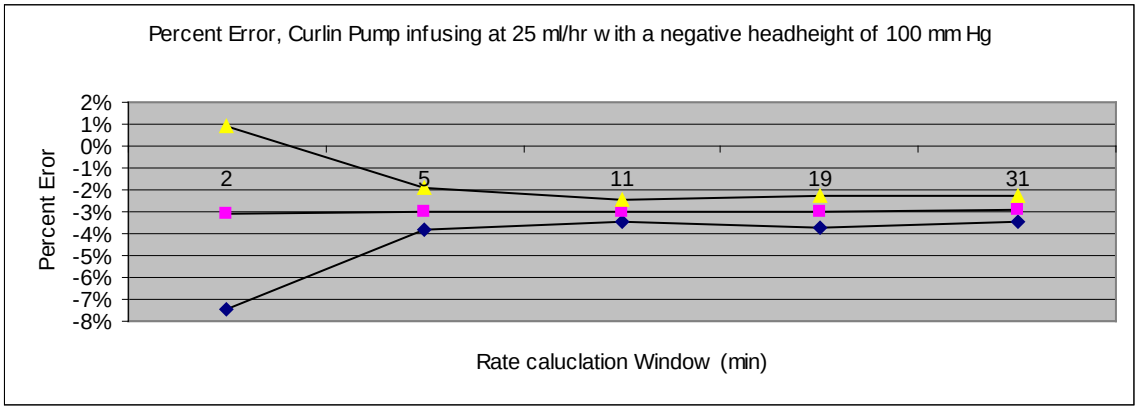
The following specifications apply to the **Curlin Infusion** Ambulatory Infusion Pump:

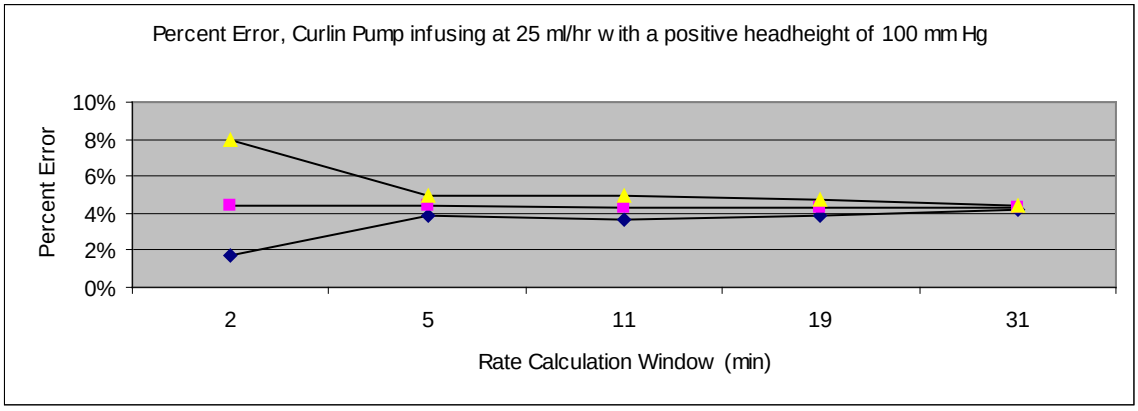
Note: Flow rates and bolus dose limits may be less than indicated in these tables based on specific customizations.

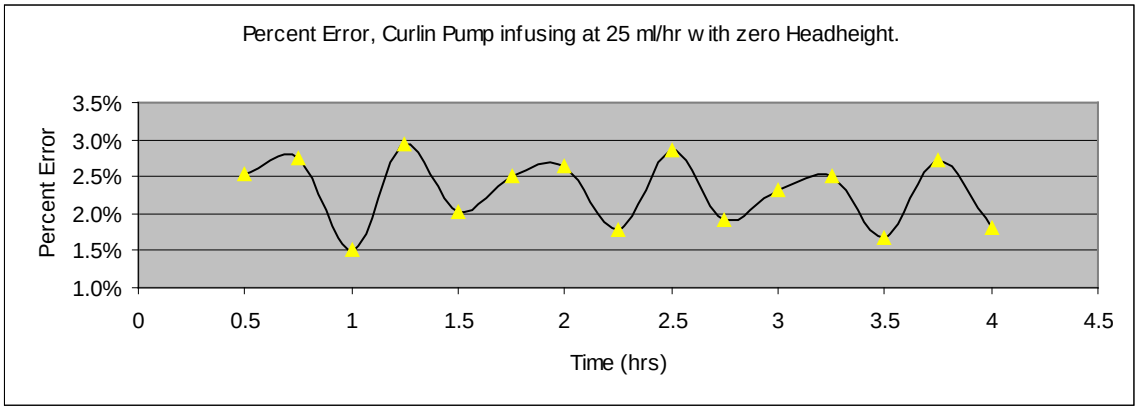
ITEM	SPECIFICATION
Pumping Mechanism:	Curvilinear peristaltic finger action
Pump Size:	
Height:	5 inches
Width:	4 inches
Depth:	2.5 inches
Weight:	18.1 ounces
Case Material:	UL rated flame-retardant, non-toxic high impact plastic
Operating Environment:	
Operating Temperature:	± 5% Accuracy: 55° to 90°F ±10% Accuracy: 45° to 55°F ±10% Accuracy: 90° to 104°F
Operating Relative Humidity:	15 to 95% non condensing
Storage Environment:	
Storage Temperature Range:	-20° to 140°F
Storage Relative Humidity:	5 to 95% non-condensing

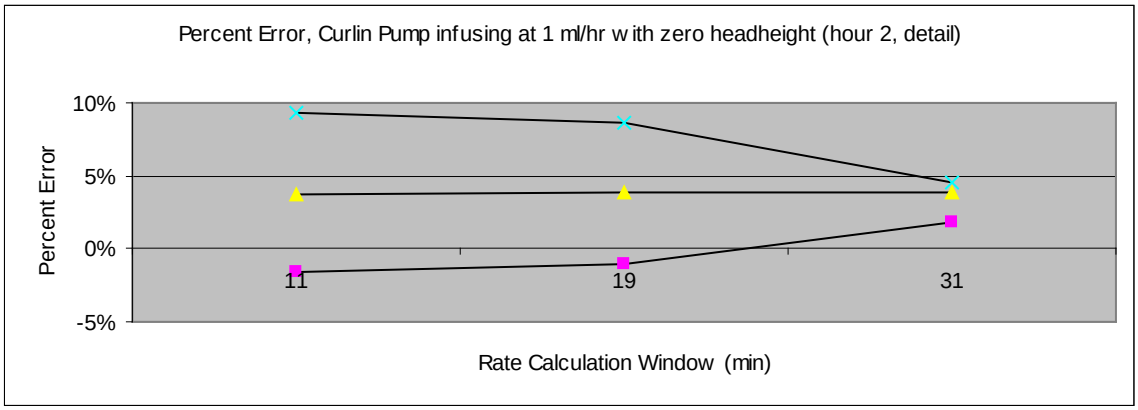
Power Requirements:	2 "C size" Single-Use Batteries AC Power
ITEM	**SPECIFICATION**
Operating Time with Full Charge Batteries:	Nominal 85 hours at a rate of 2 ml/hr
Internal Memory Battery	5 years life
Display	Graphic LCD, 4 text fields, 16 characters per field
Flow Rates	0.1 to 50 ml/hr
Bolus Rate	Up to 125 ml/hr
Volume Accuracy	± 5% for volume delivery >1 ml under the following conditions: • Temperature 55° to 90° F • Back pressure to 300 mmHg (5.8 PSI) ±10% for volume delivery >1 ml under the following conditions: • Temperature 45° to 55°F • Temperature 90° to 104°F • Back pressure to 750 mmHg (14.5 PSI) The maximum volume that may be infused under Single Fault Conditions is 0.8 ml. Recommended administration set usage not to exceed 72* hours. *follow standards of practice for set changes
Administration Sets	Proprietary **Curlin Infusion** non-DEHP, PVC administration sets available in different configurations.
Fill Volume for Administration Sets	Nonfiltered microbore sets ± 4 ml. See individual administration set label for specific fill volume
Up Occlusion Detection Time Microbore tubing with occlusion below spike	Rate — Time to Alarm 0.1 ml/hr — <20 minutes 1 ml/hr — < 4 minutes 10 ml/hr — <20 seconds 50 ml/hr — <10 seconds

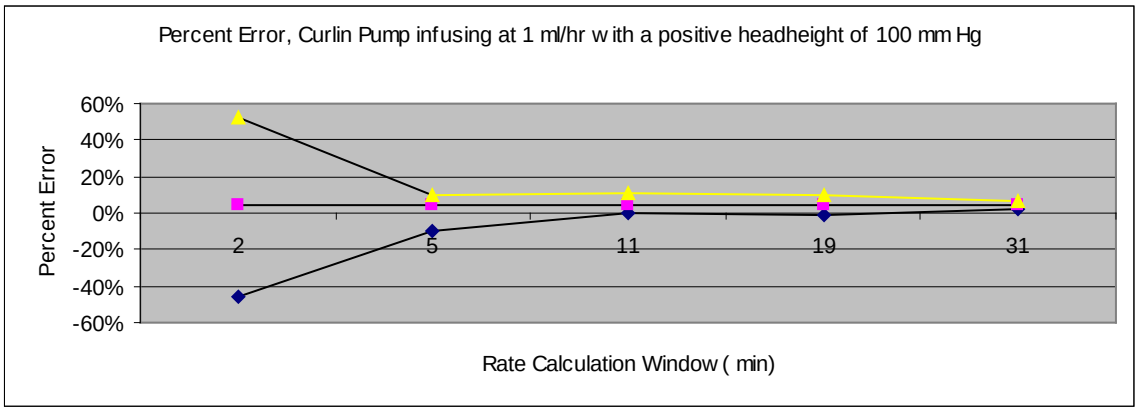
Down Occlusion Detection Time Microbore tubing with occlusion at distal Luer	Rate	Low Pressure	High Pressure
	0.1 ml/hr	<45 min	<1 hr, 30 min
	1 ml/hr	<5 min	<7 min
	10 ml/hr	<20 seconds	<45 seconds
	50 ml/hr	< 5 seconds	<15 seconds
	125 ml/hr	< 3 seconds	< 5 seconds
ITEM	**SPECIFICATION**		
Programmable Time (HH:MM) PCA	No time limit		
Bolus Volumes Microbore tubing with occlusion at distal Luer	Low Pressure: <0.1 ml High Pressure: <0.1 ml		

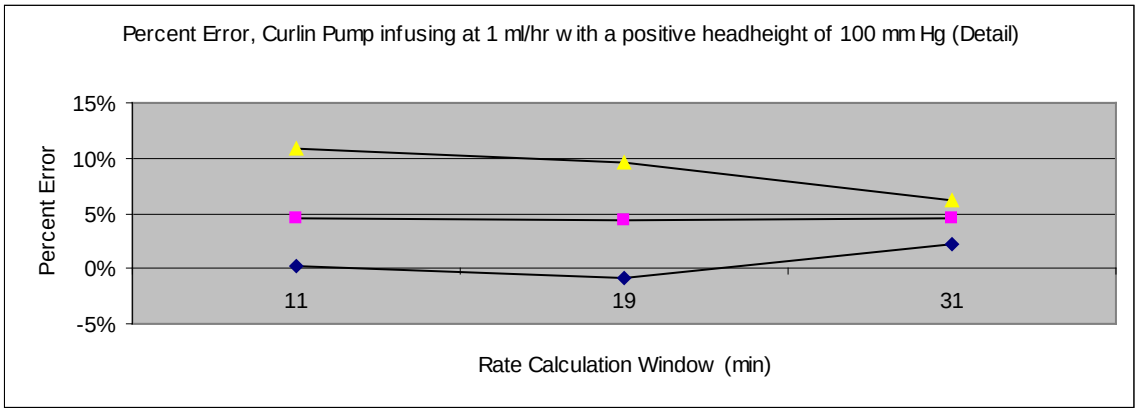


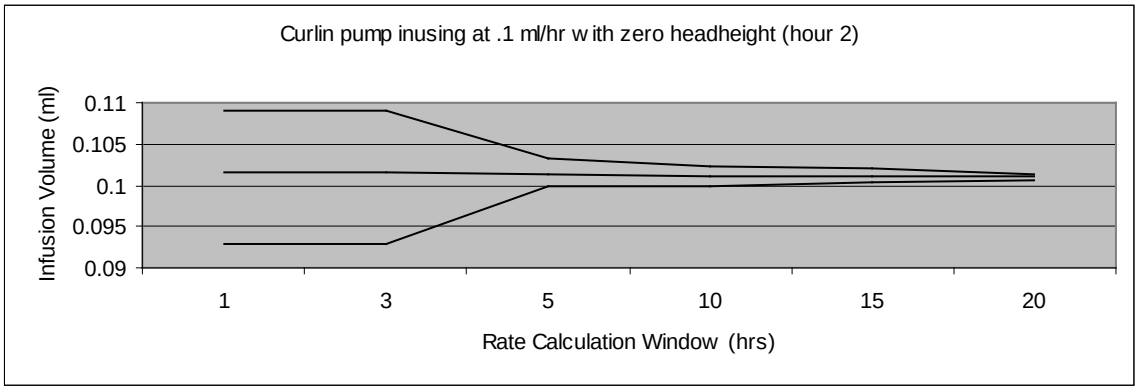


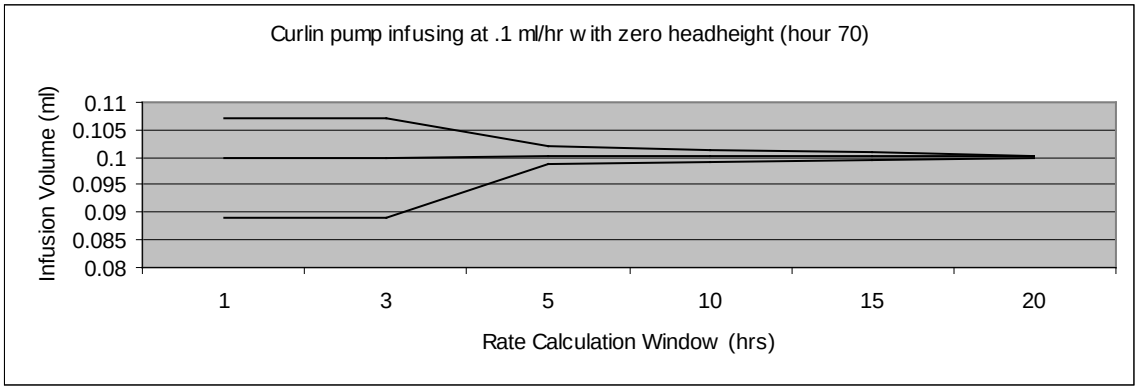

Percent Error, Curlin Pump infusing at 25 ml/hr with zero Headheight.
Percent Error
3.5%
3.0%
2.5%
2.0%
1.5%
1.0%
0
0.5
1
1.5
2
2.5
3
3.5
4
4.5
Time (hrs)

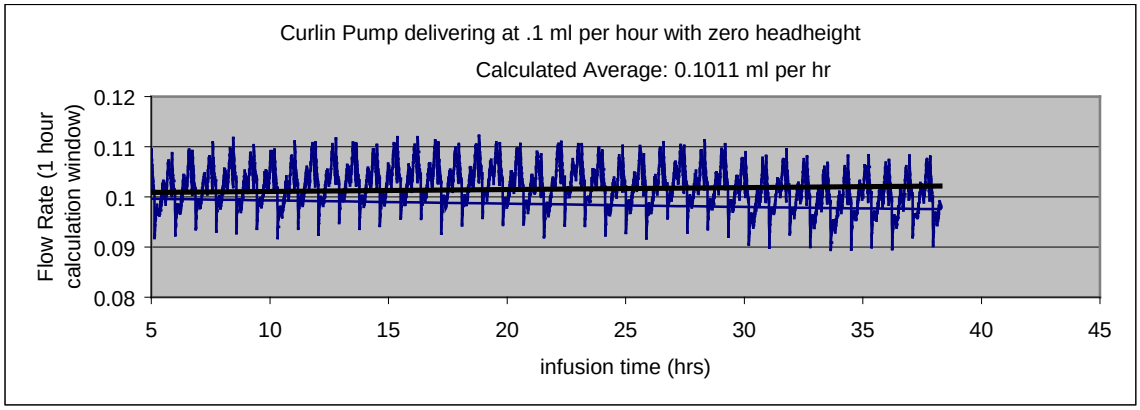

Percent Error, Curlin Pump infusing at 1 ml/hr with zero headheight (hour 2, detail)
Percent Error
10%
5%
0%
-5%
11
19
31
Rate Calculation Window (min)

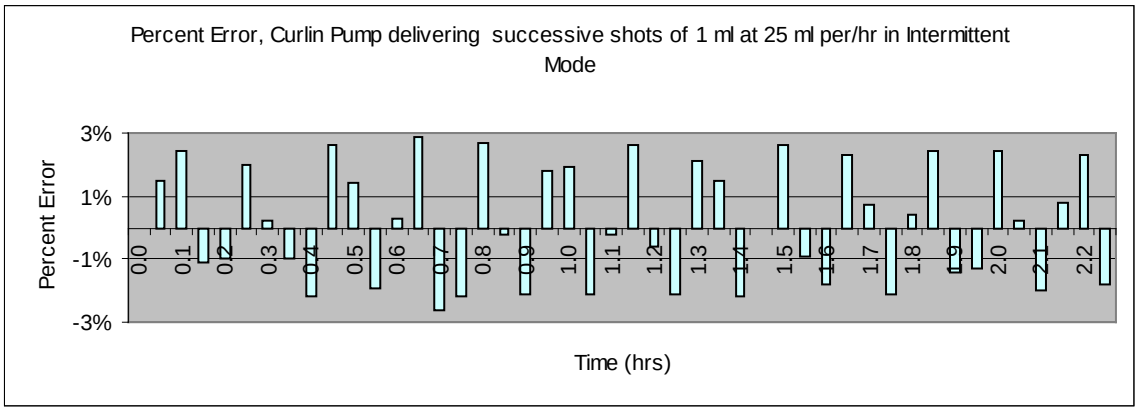

Percent Error, Curlin Pump infusing at 1 ml/hr with a positive headheight of 100 mm Hg
Percent Error
60%
40%
20%
0%
-20%
-40%
-60%
2
5
11
19
31
Rate Calculation Window (min)

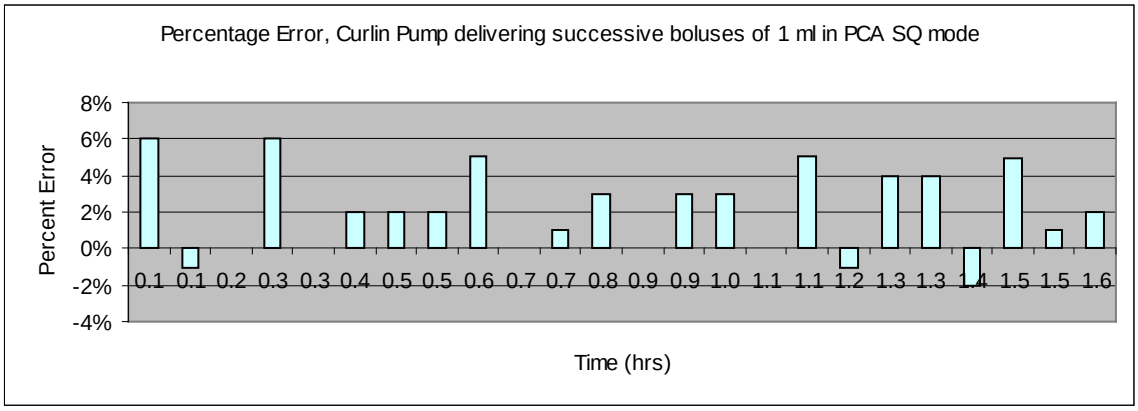

Percent Error, Curlin Pump infusing at 1 ml/hr with a positive headheight of 100 mm Hg (Detail)
Percent Error
15%
10%
5%
0%
-5%
11
19
31
Rate Calculation Window (min)

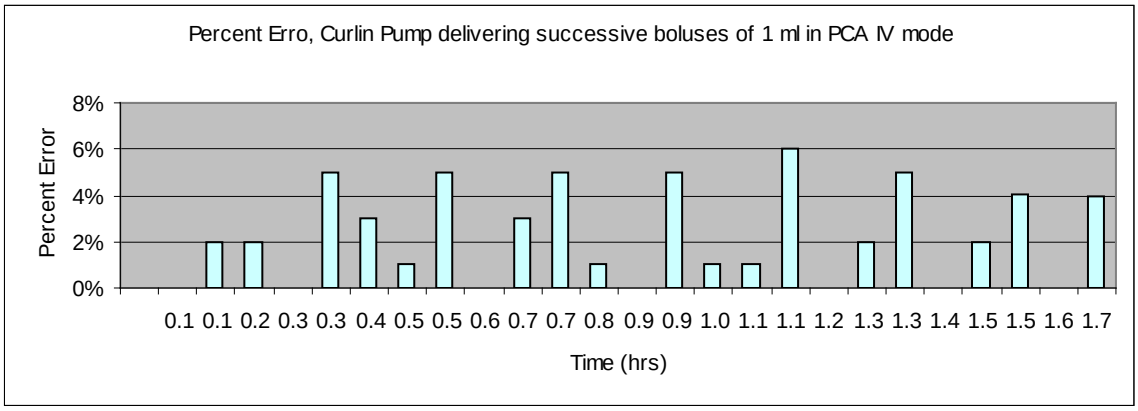

Curlin pump inusing at .1 ml/hr with zero headheight (hour 2)
Infusion Volume (ml)
0.11
0.105
0.1
0.095
0.09
1
3
5
10
15
20
Rate Calculation Window (hrs)

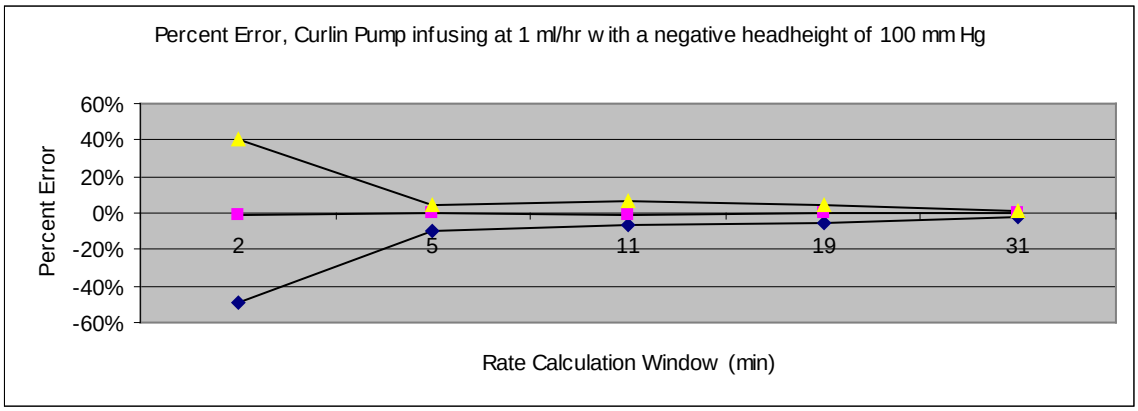

Curlin pump infusing at .1 ml/hr with zero headheight (hour 70)
Infusion Volume (ml)
0.11
0.105
0.1
0.095
0.09
0.085
0.08
1
3
5
10
15
20
Rate Calculation Window (hrs)

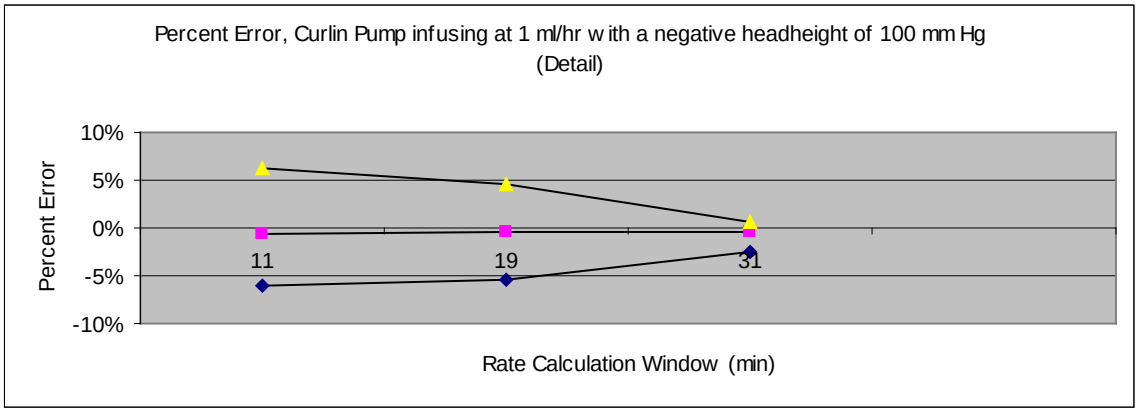

Curlin Pump delivering at .1 ml per hour with zero headheight
Calculated Average: 0.1011 ml per hr
Flow Rate (1 hour calculation window)
0.12
0.11
0.1
0.09
0.08
5
10
15
20
25
30
35
40
45
infusion time (hrs)

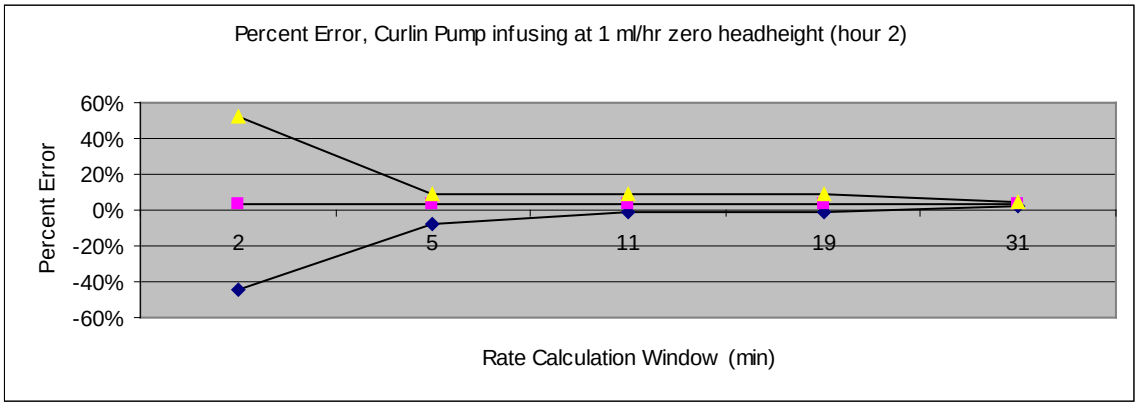

Percent Error, Curlin Pump delivering successive shots of 1 ml at 25 ml per/hr in Intermittent Mode
Percent Error
3%
1%
-1%
-3%
0.0
0.1
0.2
0.3
0.4
0.5
0.6
0.7
0.8
0.9
1.0
1.1
1.2
1.3
1.4
1.5
1.6
1.7
1.8
1.9
2.0
2.1
2.2
Time (hrs)

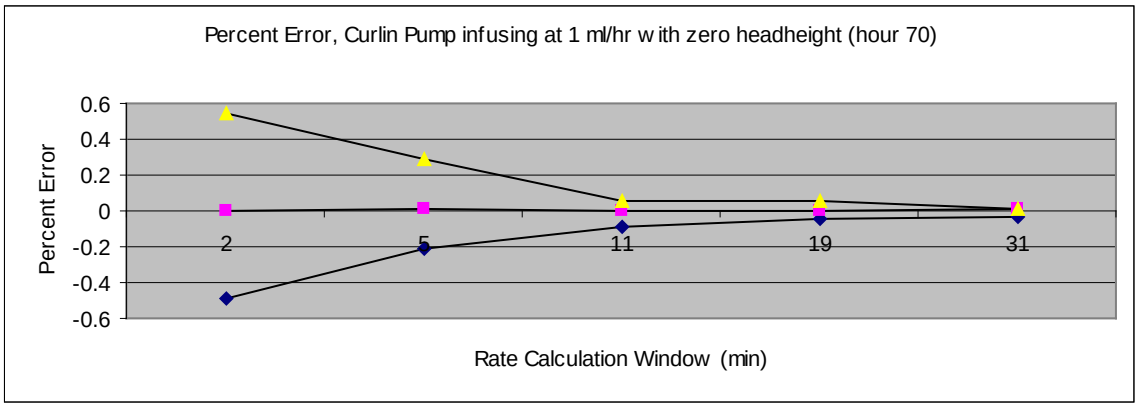

Percentage Error, Curlin Pump delivering successive boluses of 1 ml in PCA SQ mode
Percent Error
8%
6%
4%
2%
0%
-2%
-4%
0.1 0.1 0.2 0.3 0.3 0.4 0.5 0.5 0.6 0.7 0.7 0.8 0.9 0.9 1.0 1.1 1.1 1.2 1.3 1.3 1.4 1.5 1.5 1.6
Time (hrs)

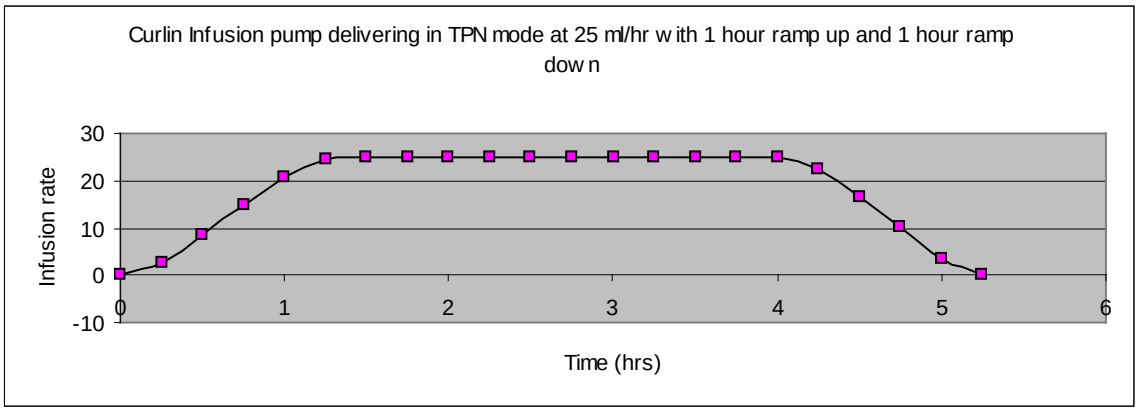

Percent Erro, Curlin Pump delivering successive boluses of 1 ml in PCA IV mode
Percent Error
8%
6%
4%
2%
0%
0.1 0.1 0.2 0.3 0.3 0.4 0.5 0.5 0.6 0.7 0.7 0.8 0.9 0.9 1.0 1.1 1.1 1.2 1.3 1.3 1.4 1.5 1.5 1.6 1.7
Time (hrs)

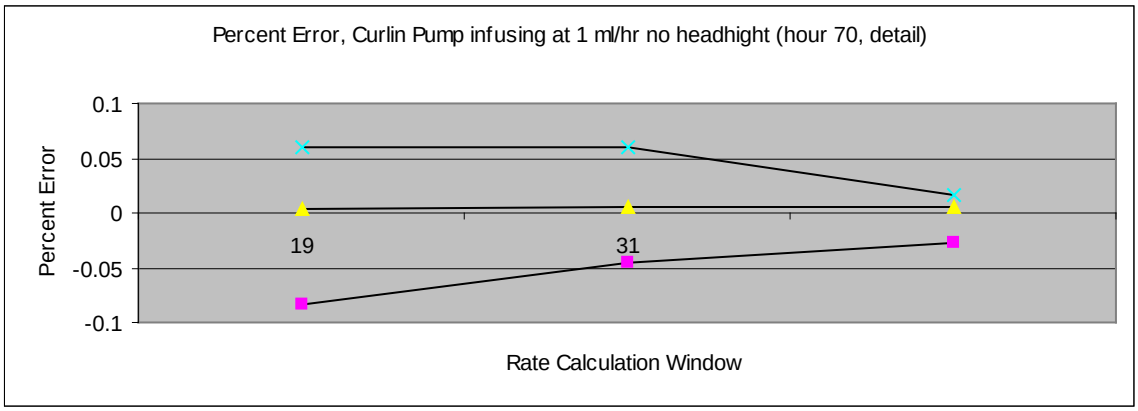

Percent Error, Curlin Pump infusing at 1 ml/hr with a negative headheight of 100 mm Hg
Percent Error
60%
40%
20%
0%
-20%
-40%
-60%
2
5
11
19
31
Rate Calculation Window (min)

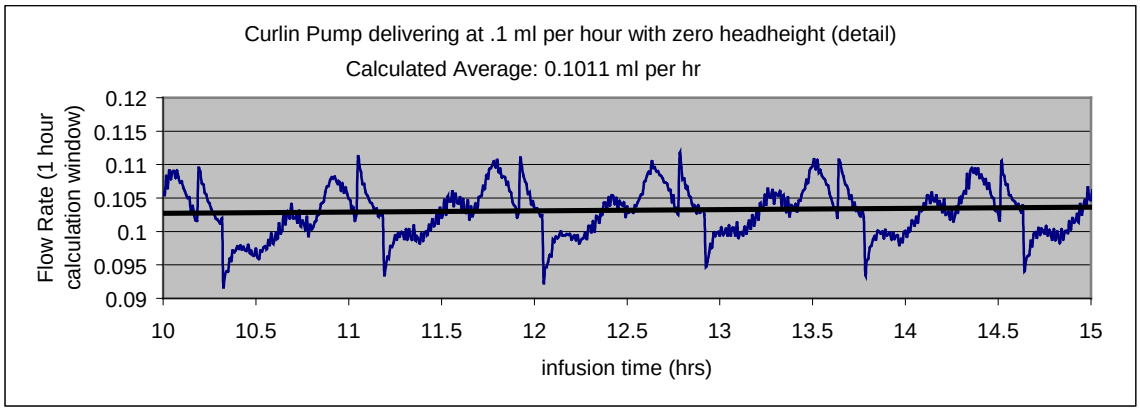

Percent Error, Curlin Pump infusing at 1 ml/hr with a negative headheight of 100 mm Hg
(Detail)
Percent Error
10%
5%
0%
-5%
-10%
11
19
31
Rate Calculation Window (min)

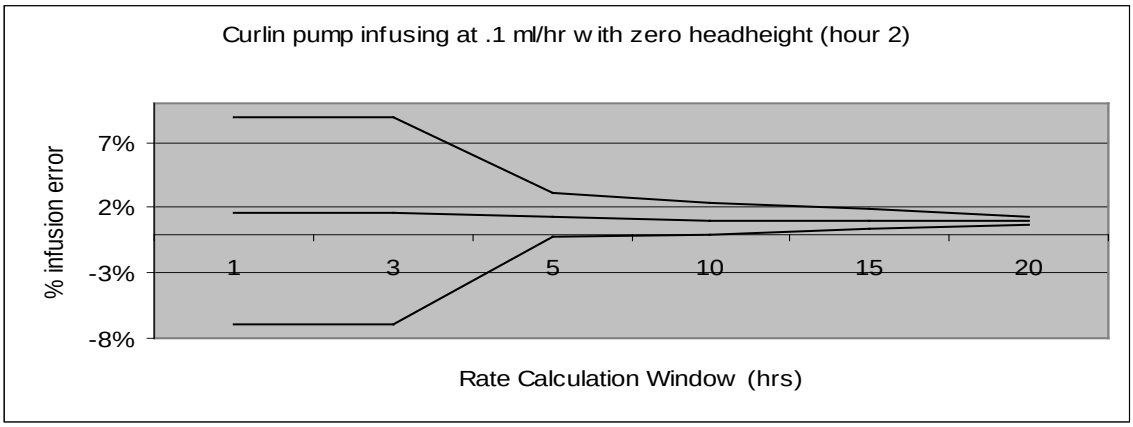

Percent Error, Curlin Pump infusing at 1 ml/hr zero headheight (hour 2)
60%
40%
20%
0%
-20%
-40%
-60%
Percent Error
2
5
11
19
31
Rate Calculation Window (min)

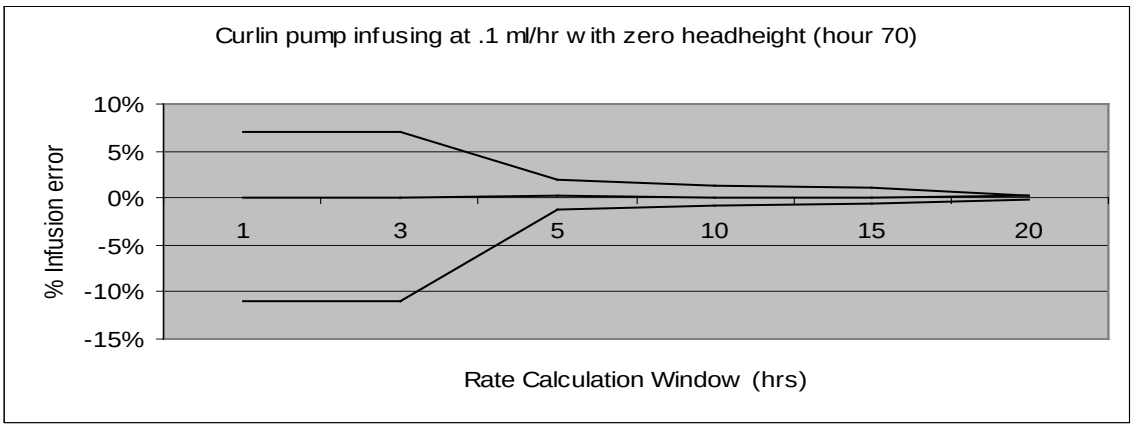

Percent Error, Curlin Pump infusing at 1 ml/hr with zero headheight (hour 70)
0.6
0.4
0.2
0
-0.2
-0.4
-0.6
Percent Error
2
5
11
19
31
Rate Calculation Window (min)


Curlin Infusion pump delivering in TPN mode at 25 ml/hr with 1 hour ramp up and 1 hour ramp down
30
20
10
0
-10
Infusion rate
0
1
2
3
4
5
6
Time (hrs)


Percent Error, Curlin Pump infusing at 1 ml/hr no headhight (hour 70, detail)
0.1
0.05
0
-0.05
-0.1
Percent Error
19
31
Rate Calculation Window


Curlin Pump delivering at .1 ml per hour with zero headheight (detail)
Calculated Average: 0.1011 ml per hr
Flow Rate (1 hour calculation window)
0.12
0.115
0.11
0.105
0.1
0.095
0.09
10
10.5
11
11.5
12
12.5
13
13.5
14
14.5
15
infusion time (hrs)


Curlin pump infusing at .1 ml/hr with zero headheight (hour 2)
% infusion error
7%
2%
-3%
-8%
1
3
5
10
15
20
Rate Calculation Window (hrs)


Curlin pump infusing at .1 ml/hr with zero headheight (hour 70)
% Infusion error
10%
5%
0%
-5%
-10%
-15%
1
3
5
10
15
20
Rate Calculation Window (hrs)

Lock Setting Table

The following table illustrates which features are controlled under which lock settings:

Pump Operation	Active Therapy	Lock OFF	Lock 1	Lock 2	Lock 3
Change Lock Settings	PCA	All settings with correct access code			
Turn Pump Off/On	PCA	Yes	Yes	Yes	Yes
Pause/Run Therapy	PCA	Yes	Yes	Yes	Yes
Access Help/Options	PCA	Yes	Yes	Yes	Yes
Check Power Source	PCA	Yes	Yes	Yes	Yes
Resume Therapy	PCA	Yes	Yes	Yes	Yes
View Rx Screens at "No to Review"	PCA	Yes	Yes	Yes	Yes
Repeat Therapy	PCA	Yes	Yes	Yes	No
Prime Pump	PCA	Yes	Yes	Yes*	No
Change Audio Level	PCA	Yes	Yes	Yes	No
Change Rx screen information	PCA	Yes	Yes	No	No
Change Rx screen at "No to Review"	PCA	Yes	Yes	No	No
Access New Program	PCA	Yes	No	No	No
Change Air in Line Setting	PCA	Yes	No	No	No
Change Down Pressure	PCA	Yes	No	No	No
Change Units (New Program)	PCA	Yes	No	No	No
Change Concentration (New Program)	PCA	Yes	No	No	No
Utilize medLIMITS Option	PCA	Yes	Yes	No	No
Program medLIMITS	PCA	Yes	No	No	No
Change Administration Route (New Program)	PCA	Yes	No	No	No
Program Loading Dose	PCA	Yes	No	No	No
Program Clinician Dose	PCA	Yes	No	No	No

Whenever a lock setting is placed in the pump it is retained unless changed by using either the BIOMED SETUP function or the OPTIONS function to change the setting. If a locked-out function is requested, such as PRIME in lock setting 3, a screen will appear requesting the entry of the access code. When the correct code is entered, the locked out function can be performed by the clinician.

Note: The lock setting remains the same, but the pump recognizes the access code, and authorizes the function to be performed without the need to change the lock setting.

* with software revision of 6R9 and higher, Prime Pump is Lock Level 2 protected.

Chapter

12

PainSmart™ IOD Catalog Numbers

PAINSMART™ IOD CATALOG NUMBERS					
Catalog Number	Units	Default Route	Fixed Route	Lock Code	Lens Color
360-1300-E1	ml		EPI	5	Yellow
360-1300-E2	ml		EPI	9	Yellow
360-1300-E3	ml	IV		9	White
360-1300-E4	ml	EPI		9	White
360-1300-E5	mg	IV		5	White
360-1300-E6	mg	IV		9	White
360-1300-E7	ml	IV		5	White
360-1300-E8	mg		IV	9	Green
360-1300-E9	ml	EPI		5	White
360-1300-E10	mg		IV	5	Green

Pump has been factory configured to specifications indicated in the above columns. Therefore, all options indicated in the manual will not be available. Pump will function according to set specifications.