# Operator’s Manual

## English

### TABLE OF CONTENTS

<table>
<thead>
<tr>
<th>Section</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>INTENDED USE</td>
<td>2</td>
</tr>
<tr>
<td>SUMMARY AND EXPLANATION</td>
<td>2</td>
</tr>
<tr>
<td>PRINCIPLES OF OPERATION</td>
<td>3</td>
</tr>
<tr>
<td>DESCRIPTION</td>
<td>5</td>
</tr>
<tr>
<td>IMPORTANT LABELS AND SYMBOLS</td>
<td>10</td>
</tr>
<tr>
<td>SPECIFICATIONS</td>
<td>12</td>
</tr>
<tr>
<td>PREPARING THE INSTRUMENT</td>
<td>13</td>
</tr>
<tr>
<td>BATTERY PRECAUTIONS</td>
<td>13</td>
</tr>
<tr>
<td>STARTING THE INSTRUMENT</td>
<td>25</td>
</tr>
<tr>
<td>RUNNING A TEST</td>
<td>26</td>
</tr>
<tr>
<td>QUALITY CONTROL (QC)</td>
<td>35</td>
</tr>
<tr>
<td>OPERATING PRECAUTIONS</td>
<td>37</td>
</tr>
<tr>
<td>LIMITATIONS</td>
<td>38</td>
</tr>
<tr>
<td>RESULTS MANAGEMENT</td>
<td>38</td>
</tr>
<tr>
<td>COMMUNICATION WITH A PC OR NETWORK</td>
<td>44</td>
</tr>
<tr>
<td>CONFIGURATION MANAGER</td>
<td>46</td>
</tr>
<tr>
<td>MASTER FILE MAINTENANCE</td>
<td>66</td>
</tr>
<tr>
<td>CONFIGURATION MANAGER UTILITIES</td>
<td>67</td>
</tr>
<tr>
<td>TROUBLESHOOTING</td>
<td>72</td>
</tr>
<tr>
<td>SERVICE AND MAINTENANCE</td>
<td>76</td>
</tr>
<tr>
<td>SAFETY STANDARDS</td>
<td>77</td>
</tr>
<tr>
<td>INDEX</td>
<td>79</td>
</tr>
</tbody>
</table>

This manual is published by Accriva Diagnostics, Inc. (Accriva) for use with the Hemochron Signature Elite Whole Blood Microcoagulation System.

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**INTENDED USE**

The Hemochron Signature Elite® Whole Blood Microcoagulation System is a battery-operated, hand-held instrument that performs individual point-of-care coagulation tests on fresh or citrated whole blood. These tests include: Activated Clotting Time (ACT+ and ACT-LR), Activated Partial Thromboplastin Time (APTT and APTT Citrate), and Prothrombin Time (PT and PT Citrate). The system is intended to be used only with test cuvettes that are available from the manufacturer.

Data management capabilities are included with the instrument. These capabilities include storage of up to 600 patient results and 600 quality control results, designation of quality control levels, tagging of test results with date and time, entry of Patient ID and/or Operator ID or Operator PIN, and printing of results.

Hemochron® Configuration Manager software is included. This software allows the user to connect a personal computer to the instrument and perform system configuration functions using the fast and convenient Microsoft® Windows® user interface. Hemochron ReportMaker™ software, provided separately, allows the user to connect a personal computer to an instrument and perform various data management and data reporting functions.

For *in vitro* Diagnostic Use. For professional use. Rx only.

**SUMMARY AND EXPLANATION**

Events that lead to formation of a blood clot are simplified in coagulation theory into two interactive coagulation cascades that are referred to as the *intrinsic* and the *extrinsic* pathways.

The clotting factors involved in these pathways are numbered I through V and VII through XIII. The intrinsic pathway begins with the contact activation of factor XII and, through the interaction of several coagulation factors, results in the conversion of soluble fibrinogen to insoluble fibrin strands. The extrinsic pathway is initiated through the interaction of tissue factor with factor VII. Platelets, essential co-factors in this reaction, provide the platelet phospholipid surface on which coagulation reactions occur.

Activated Clotting Time (ACT+ and ACT-LR), Activated Partial Thromboplastin Time (APTT and APTT Citrate), and Prothrombin Time (PT and PT Citrate) tests are general coagulation screening tests that are used to measure the functionality of the blood coagulation cascade. The ACT test is the method of choice for monitoring heparin therapy during cardiac surgery and PCI. Fresh whole blood is added to an activator (Celite®, silica, kaolin, or glass particles), and then timed for the formation of a clot. The type of activator used will affect the degree of prolongation of the ACT to a given heparin dose.

The Hemochron®. ACT+ test uses a mixture of silica, kaolin, and phospholipids as an activator to create a rapid and highly sensitive alternative to existing ACT tests. This test demonstrates linearity at heparin concentrations ranging from 1.0 to 6.0 units of heparin per mL of blood, and it is not affected by high dose aprotinin therapy.

The Hemochron®. ACT-LR test uses a Celite activator due to its excellent heparin sensitivity. The test demonstrates linearity at heparin concentrations up to 2.5 units of heparin per mL of blood. The test is not intended for use with aprotinin therapy.

The Hemochron®. APTT test measures the intrinsic coagulation pathway and involves all coagulation factors except factors VII and III (tissue factor). The APTT test improves the PTT test through use of a contact activating substance, which standardizes activation of Factor XII to provide a more precise and sensitive assay for low level heparin monitoring. The Hemochron®. APTT test is formulated to provide optimal heparin sensitivity at heparin concentrations up to 1.5 units of heparin per mL of blood.

The Hemochron®. APTT Citrate test performs the same measurement as the APTT test, using a citrated whole blood sample.
The Hemochron PT test measures the extrinsic coagulation pathway and is sensitive to coagulation factors VII, X, V, II, and fibrinogen. PT results may be abnormal in patients with liver disease or Vitamin K deficiency. The test is widely used to monitor oral anticoagulant therapy. The PT test is a unitized test system utilizing a highly sensitive thromboplastin for improved specificity and sensitivity.

The Hemochron PT Citrate test performs the same measurement as the PT test, using a citrated whole blood sample.

**Note:** Hemochron Configuration Manager V3.0 or higher is required and is included with the Hemochron Signature Elite instrument.

**PRINCIPLES OF OPERATION**

The Hemochron Signature Elite Whole Blood Microcoagulation System provides many features for ease of use and reliability, including a patented clot detection system, a data storage module, interfaces for a laboratory computer and/or printer, a streamlined user-interface panel, and an integrated barcode scanner.

The system measures whole blood clotting times using Hemochron disposable single-use cuvettes. Each cuvette contains all of the reagents necessary for a specified test.

The operator inserts a cuvette for the test into the instrument and then (if desired) enters information about the sample or scans the information from a barcode using the integral barcode scanner. After the cuvette has warmed to 37 °C ±1.0 °C, the instrument beeps, signaling the operator that a blood sample can be added to the cuvette and the test started.

The operator then places a drop of blood in the sample well of the cuvette and presses the **START** key. The instrument measures the required volume of blood and automatically moves it into the cuvette test channel, where it is mixed with reagents. The remainder of the blood sample, not needed for testing, is automatically drawn out of the sample well and into an enclosed waste channel on the cuvette.

After mixing with the reagent, the sample is moved back and forth at a predetermined rate within the test channel and monitored for clot formation. The test channel is maintained at 37 °C ±1.0 °C during the test.

The rate of movement of the sample is monitored by a series of LED optical detectors that are aligned with the test channel. When the blood clots, the flow of the blood sample within the test channel is impeded, reducing its rate of flow between the optical detectors. This reduction in flow below a predetermined value signals to the instrument that a clot has formed. The instrument also emits an audible beep when clot formation occurs, indicating the end of the test. An internal timer measures the elapsed time between the start of the test and the clot formation. During the test, the whole blood clotting time (in seconds) is displayed.

APTT and APTT Citrate results are displayed as plasma equivalent (PE) values, PT and PT Citrate results are displayed as the International Normalized Ratios (INR) and PE values, and ACT+ and ACT-LR results are displayed as Celite ACT equivalent time.

The results will remain on the display while the cuvette remains in the instrument. The result can be automatically printed along with the time and date the test was run, the Patient ID, Operator ID, and other information, if entered. The result is also saved in an internal database. Up to 600 patient test results and 600 quality control test results can be stored in the instrument for later printing or downloading.

Individual Hemochron Signature Elite instruments can be customized so that designated quality control tests must be performed whenever a specified period of time has elapsed and so that information for cuvettes and liquid controls can be included in test records. The instruments can also be configured so that only authorized operators can operate the system and/or operators cannot perform certain functions such as deleting test results from the instrument database. These and other configuration options are entered using Hemochron Configuration Manager Software on a personal computer.
Definitions and Terms
The following acronyms and abbreviations are used in this manual, instrument screens, and printouts:

- **ACT**: Activated Clotting Time
- **APTT**: Activated Partial Thromboplastin Time
- **CVlot**: Cuvette Lot Number
- **DB**: Database
- **EQC**: Electronic Quality Control (System Verification)
- **HCM**: Hemochron Configuration Manager
- **INR**: International Normalized Ratio
- **idms**: Integrated Data Management System
- **LQC**: Liquid Quality Control
- **OID**: Operator Identification Number
- **PID**: Patient Identification Number
- **PIN**: Operator Personal Identification Number
- **POCC**: Point of Care Coordinator
- **PT**: Prothrombin Time
- **QC**: Quality Control
- **QClot**: Quality Control Lot Number
- **TQC**: Temperature Quality Control
DESCRIPTION

The Hemochron Signature Elite instrument is a hand-held device. It contains a test chamber which warms a test cuvette to the required temperature, and it performs all operations to measure the clotting time of a whole blood sample after it is placed in the test cuvette and the test is started by the operator.

Operator instructions are shown on the display panel, and the operator enters commands and information using the keypad. When the test is completed, the results are shown on the display panel and stored in system memory for printing or downloading.

The instrument includes a barcode scanner for reading of barcode labels.

Keypad

The front panel contains a keypad with various action keys as well as a number pad. The operator uses the keypad to select a command or enter information.

Action keys display a menu of commands according to the label on the key. For example, the PRINT/SCAN key displays printing commands and enables scanning. Number pad (or option) keys are used to enter characters for IDs or numbers for selection of commands.

Note: The 0 (zero) key is used as both a numeral key and an action key.

<table>
<thead>
<tr>
<th>Key</th>
<th>Purpose</th>
</tr>
</thead>
<tbody>
<tr>
<td>START</td>
<td>Turn on the instrument or turn off instrument.</td>
</tr>
<tr>
<td></td>
<td>Initiate a test after the test cuvette has reached 37 °C ±1.0 °C and the sample is added.</td>
</tr>
</tbody>
</table>
PRINT/SCAN
Print results on an external printer for the last test result, patient database, quality control database, or all test results for a specified patient.
Test the printer.
Enter a customized header.
Print instrument system information.
Scan a barcode ID when Enter PID, Enter OID, Enter PIN, Enter CVLot, Enter QCLot, or Scan Lot is displayed.

NOTE
Select a laboratory-defined note (from nine choices programmed via Hemochron Configuration Manager) for entry into the record for the current test.
Toggle between alphabetic (ABC) and numeric (123) characters when Enter PID, Enter OID, Enter PIN, Enter CVLot, or Enter QCLot is displayed.

ID
Enter a Patient ID or Operator ID.

QC
Specify that the test being run is a normal (level 1) control or an abnormal (level 2) control.
(For QC Lockouts) Display the length of time remaining before Quality Control testing for a particular test must be successfully completed.
Initiate an EQC (Electronic Quality Control) test.

DATABASE
Display the number of patient or QC test results that are currently stored in the instrument and the number of additional test results that can be stored.
Display results of the stored patient or QC tests.
Delete patient or QC records.
Initiate POCT-1A communication.

0 through 9
Select a menu option.
Enter information (e.g., a Patient ID, an Operator ID, cuvette lot number, QC lot number, a time, or a date).
Display the main menu by pressing the Zero key when no tests are running and no menu is active.

CANCEL
Cancel an operation and return to the previous state.

ENTER
Accept an entry (e.g., a Patient ID, an Operator ID, a time, or a date).

Display Panel
The instrument communicates with the operator by means of the display panel. For example, Add Sample and Press Start are alternately displayed after a test cuvette reaches temperature:

![Display Panel Example](image)

The display is illuminated to enhance visibility in low light conditions. To conserve power during battery operation, the display is not illuminated when the test well is empty or when a test is running. The display is momentarily illuminated after a key is pressed or when results of a completed test are displayed. While a test is running, the display can be momentarily illuminated by pressing the CANCEL key, the ENTER key, or any numeric key.
The action keys are used to display and activate various commands for operating and configuring the system. The principal menus are shown below:

<table>
<thead>
<tr>
<th>Menu</th>
<th>Commands</th>
<th>Access</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Main Menu</strong></td>
<td><strong>MAIN MENU</strong></td>
<td>Press the 0 (zero) key when no tests are running and no other menu is active.</td>
</tr>
<tr>
<td></td>
<td>1-Power Status</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-System Time</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-System Date</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4-Shut Down</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5-System Info</td>
<td></td>
</tr>
<tr>
<td></td>
<td>6-Configuration</td>
<td></td>
</tr>
<tr>
<td>**Configuration **</td>
<td><strong>CONFIGURATION</strong></td>
<td>Press the 6 key while the Main Menu is displayed.</td>
</tr>
<tr>
<td></td>
<td>1-Contrast</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-Brightness</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-Flashlight</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4-Language Menu</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5-Supervisor</td>
<td></td>
</tr>
<tr>
<td><strong>Supervisor</strong></td>
<td><strong>SUPERVISOR</strong></td>
<td>Press the 5 key while the Configuration Menu is displayed, enter the supervisor PIN at the prompt, then press and hold the Enter key.</td>
</tr>
<tr>
<td></td>
<td>1-Scan OID</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-System Time</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-Supervisor PIN</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4-Time Zone</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5-Print Options</td>
<td></td>
</tr>
<tr>
<td></td>
<td>6-Communications</td>
<td></td>
</tr>
<tr>
<td><strong>Print Options</strong></td>
<td><strong>PRINT OPTIONS</strong></td>
<td>Press the 6 key while the Supervisor Menu is displayed.</td>
</tr>
<tr>
<td></td>
<td>1-Print Operator</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-Print CUU Lots</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-Print QC Lots</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4-Baudrate</td>
<td></td>
</tr>
<tr>
<td><strong>Communications</strong></td>
<td><strong>COMMUNICATIONS</strong></td>
<td>Press the 7 key while the Supervisor Menu is displayed.</td>
</tr>
<tr>
<td></td>
<td>1-POCT</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-Port</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-Push POCT OFF</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4-XPORT Setup</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5-XPORT COM/LSET</td>
<td></td>
</tr>
<tr>
<td></td>
<td>6-ITC &gt;&gt; COM</td>
<td></td>
</tr>
<tr>
<td><strong>QC Selects Menu</strong></td>
<td><strong>QC SELECTS</strong></td>
<td>Press the QC key while a cuvette is inserted in the instrument.</td>
</tr>
<tr>
<td></td>
<td>1-QC Normal</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-QC Abnormal</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-Patient Result</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4-QC Status</td>
<td></td>
</tr>
<tr>
<td><strong>QC Status Menu</strong></td>
<td><strong>QC Status</strong></td>
<td>Press the QC key before a cuvette is inserted in the instrument (or, select 4 from the QC Selects menu.)</td>
</tr>
<tr>
<td></td>
<td>Press 1-Run EQC</td>
<td></td>
</tr>
<tr>
<td></td>
<td>APTT 10:00</td>
<td></td>
</tr>
<tr>
<td></td>
<td>ACTT 15:30</td>
<td></td>
</tr>
<tr>
<td></td>
<td>PT 18:20</td>
<td></td>
</tr>
<tr>
<td></td>
<td>ACT+ 20:30</td>
<td></td>
</tr>
<tr>
<td></td>
<td>9-Next Page</td>
<td></td>
</tr>
<tr>
<td><strong>Notes Menu</strong></td>
<td><strong>Notes</strong></td>
<td>Press the NOTE key after a cuvette is inserted in the instrument.</td>
</tr>
<tr>
<td></td>
<td>1-9-note 0-erase</td>
<td></td>
</tr>
<tr>
<td></td>
<td>ENTER-toggle</td>
<td></td>
</tr>
<tr>
<td></td>
<td>CANCEL to Exit</td>
<td></td>
</tr>
<tr>
<td></td>
<td>No Note</td>
<td></td>
</tr>
<tr>
<td></td>
<td>No Note</td>
<td></td>
</tr>
<tr>
<td><strong>ID Selects Menu</strong></td>
<td><strong>ID SELECTS</strong></td>
<td>Press the ID key while a cuvette is inserted in the instrument.</td>
</tr>
<tr>
<td></td>
<td>1-Enter PID</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-Enter OID</td>
<td></td>
</tr>
<tr>
<td><strong>Print/Scan</strong></td>
<td><strong>PRINT MENU-1</strong></td>
<td>Press the PRINT/SCAN key.</td>
</tr>
<tr>
<td>Menu 1</td>
<td>1-Print Last</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-Print PAT DB</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-Print QC DB</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4-By PAT Menu</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5-By QC Menu</td>
<td></td>
</tr>
<tr>
<td></td>
<td>6-Print Check</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-Next Page</td>
<td></td>
</tr>
<tr>
<td>Menu</td>
<td>Commands</td>
<td>Access</td>
</tr>
<tr>
<td>----------------------</td>
<td>---------------------------</td>
<td>----------------------------------------------------------------</td>
</tr>
<tr>
<td>Print by Patient Menu</td>
<td></td>
<td>Press the 4 key while the Print Menu 1 is displayed.</td>
</tr>
<tr>
<td></td>
<td>BY PAT MENU</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1-Print by PID</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-Print by OID</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-Print by Date</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4-Print by Assay</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5-Print by CVIlot</td>
<td></td>
</tr>
<tr>
<td>Print by QC Menu</td>
<td></td>
<td>Press the 5 key while the Print Menu 1 is displayed.</td>
</tr>
<tr>
<td></td>
<td>BY QC MENU</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1-Print by OID</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-Print by Date</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-Print by Assay</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4-Print by CVIlot</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5-Print by QCIClot</td>
<td></td>
</tr>
<tr>
<td>Print/Scan Menu 2</td>
<td></td>
<td>Press the 7 key while Print Menu 1 is displayed.</td>
</tr>
<tr>
<td></td>
<td>PRINT MENU-2</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1-Enter Header</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-Print System</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-Test SCAN</td>
<td></td>
</tr>
<tr>
<td>Database Menu</td>
<td></td>
<td>Press the DATABASE key.</td>
</tr>
<tr>
<td></td>
<td>DATABASE</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1-Totals</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2-ECI Erase</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3-QC Erase</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4-Display Pat</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5-Display QC</td>
<td></td>
</tr>
<tr>
<td></td>
<td>6-ITC &gt;&gt; COM</td>
<td></td>
</tr>
</tbody>
</table>

Execute a command on a menu by pressing the action key for that command. For example, if the Main menu is displayed, press the 4 key to shut down the system, or press the 5 key to display system information.

*Note: The main menu cannot be accessed if a cuvette is in the test well. If a cuvette is used to turn on the instrument, it must be removed to access the main menu.*
Test Cuvettes
Tests are performed with single-use disposable Hemochron test cuvettes. Each test cuvette contains a label, a sample well, a test channel containing reagents, an enclosed waste reservoir, and one or more optical detection windows.

The cuvette label is human- and instrument-readable. The current test is automatically read by the instrument and displayed for confirmation when the operator inserts the cuvette into the test chamber. After the cuvette has warmed, Add Sample and Press Start are displayed by the instrument, alerting the operator to place a drop of blood in the sample well and begin the test by pressing the START key.

Note: Refer to the package insert accompanying the Hemochron test cuvettes for storage and handling instructions.

Features
The Hemochron Signature Elite has a number of performance and convenience features. These are summarized below:

- the system is portable for bedside use
- fresh whole blood or citrated whole blood can be used (assay dependent)
- only one drop of blood is required
- results are available in minutes
- results are displayed appropriately as whole blood, plasma equivalent, Celite equivalent seconds, or INR
- test type is automatically read from the cuvette
- barcode or manual entry of identification numbers, cuvette lot information, and quality control lot information
- test results are automatically stamped with date and time
- results from 600 patient tests and 600 QC tests can be stored
- stored results can be printed and/or downloaded to a personal computer
- stored results can be printed by Patient ID
- instrument self checks are automatically performed
- internal Electronic Quality Control (EQC) is designed to check instrument operation at two levels
- Electronic Quality Control (EQC) levels selectable for 300 seconds or 500 seconds using Hemochron Configuration Manager software
- internal temperature verification is designed to check test chamber temperature
- the display is illuminated for viewing in low light
- the user is alerted when the battery is low
- an optional printer can be connected
- the system can be customized using a personal computer and Hemochron Configuration Manager software
- User Notes can be entered to add additional information to the test results
- Patient/QC test reports can be created using a personal computer and data management software programs
- POCT 1A compliant
- Operator certification lockout
- Cuvette expiration lockout
- QC expiration lockout
Instrument Lockouts

The instrument can be configured to allow use only by authorized operators and/or to allow use only if specified QC has been performed. If one or more of these lockouts is enabled, processing of the lockout(s) occurs before any other instrument functions.

Prewarming

If the lockout checks are passed, a cuvette is automatically prewarmed to 37 °C ±1.0 °C when it is placed in the test chamber. The instrument will sound one short beep when pre-warming is complete.

Test Timeout

Incubation of a cuvette stops automatically if **START** is not pressed within five minutes after **Add Sample** and **Press Start** are displayed.

Automatic Shutdown

The instrument will automatically shut down after being idle for five minutes if there is no cuvette inserted in the test chamber and the instrument is running on the battery.

*Note: Results that are greater than the time noted in the package insert for a test are beyond the sensitivity range of the test. They should be repeated immediately and, if confirmed, reported as greater than the maximum sensitivity range.*

IMPORTANT LABELS AND SYMBOLS

Serial Number Label

A serial number label on the rear of the Hemochron Signature Elite instrument alerts users to the following:

1. The barcode scanner contains an IEC 60825-1 class 1 laser
2. **SN** Serial Number of Device
3. **IVD** For *in vitro* Diagnostic Use
4. **!!** Attention - Read accompanying documentation or instructions
5. **/** Manufactured date
6. **/** Manufacturer
7. **/** Medical Equipment per Annex 1A, Item 8 Directive 2002/96/EC For Electronic Equipment Waste – Contact Technical Support @ 877-579-4441 (toll free)
Other Symbols

Before using the Hemochron Signature Elite system, it is essential that the contents of this Operator’s Manual, any labels on the instrument or its packaging, and instructions accompanying Hemochron cuvettes are read and understood by the operator. These materials make reference to additional symbols that are explained below:

- **Handle with care.** Handle and open the container with care.
- **Expiration Date of Cuvettes**
- **Lot Number of Cuvettes**
- **Do Not Reuse Cuvettes – Single Use Only**
- **Upper and Lower Temperature Limitations (For Storage or Use)**
- **Consult Instructions for Use**
- **Input Port for DC Power Cord from AC/DC Power Module - Polarity, VDC and A Input.**
- **RS232 Output Port for Data Transfer.**
- **Ethernet Output Port for Data Transfer.**
- **Laser Aperture Label.** Alerts the user that a class 1 or class 2 laser Radiation is emitted from the aperture. The label is located on the side of the instrument, next to the aperture.
SPECIFICATIONS

Specifications for the Hemochron Signature Elite Whole Blood Microcoagulation System are listed below.

### Dimensions and Weight

<table>
<thead>
<tr>
<th>Dimension</th>
<th>Measurement</th>
</tr>
</thead>
<tbody>
<tr>
<td>Depth</td>
<td>9.4 cm (3.7 in)</td>
</tr>
<tr>
<td>Width</td>
<td>19 cm (7.5 in)</td>
</tr>
<tr>
<td>Height</td>
<td>5 cm (2.0 in)</td>
</tr>
<tr>
<td>Weight</td>
<td>0.53 kg (1.2 lbs)</td>
</tr>
</tbody>
</table>

### Performance

- **Test Precision**: ≤ 10% C.V. for whole blood samples

### Operation

<table>
<thead>
<tr>
<th>Specification</th>
<th>Details</th>
</tr>
</thead>
<tbody>
<tr>
<td>Test Chamber</td>
<td>1</td>
</tr>
<tr>
<td>Timing Range</td>
<td>0 second to 1005 seconds</td>
</tr>
<tr>
<td>Incubation Temperature</td>
<td>37 °C ±1.0 °C</td>
</tr>
<tr>
<td>Incubation Warm-Up Time</td>
<td>30 seconds to 90 seconds</td>
</tr>
<tr>
<td>Full-Charge Operating Time</td>
<td>&gt; 2 hours</td>
</tr>
<tr>
<td>Battery Life</td>
<td>500 recharges</td>
</tr>
<tr>
<td>Battery Type</td>
<td>Lithium Ion</td>
</tr>
<tr>
<td>Throughput (Full Charge)</td>
<td>49 test cycles (at 150 sec per test)</td>
</tr>
<tr>
<td>Operating Environment</td>
<td>15 °C to 30 °C</td>
</tr>
<tr>
<td>Transport and Storage</td>
<td>-20 °C to 50 °C</td>
</tr>
</tbody>
</table>

### AC/DC Power Module

<table>
<thead>
<tr>
<th>Power Module</th>
<th>Details</th>
</tr>
</thead>
<tbody>
<tr>
<td>Input Power</td>
<td>100 to 240 VAC, 50 to 60 Hz</td>
</tr>
<tr>
<td>Output Power</td>
<td>12.0 VDC, 3.4 Amps (40 Watts)</td>
</tr>
</tbody>
</table>

### Barcode Reader

<table>
<thead>
<tr>
<th>Barcode Reader</th>
<th>Description</th>
</tr>
</thead>
</table>

**Laser Class**

IEC 60825-1 class 1 laser product complies with 21 CFR 1040.10 and 1040.11 except for deviations pursuant to Laser Notice No. 50 dated June 24, 2007.

| Laser Wavelength                    | Class 1: 630-670 nm                                                         |
|-------------------------------------|                                                                             |
| Laser Power (peak)                  | Class 1: 0.7mW @peak                                                        |

### Calibration

The Signature Elite instrument is calibrated at the manufacturing facility to test and verify all functions. The instrument is also self-calibrating, as all instrument functions are continuously monitored and verified by the instrument software when a test is performed. The instrument does not require additional calibration by the user.
PREPARING THE INSTRUMENT

Unpacking and Inspection

*Note:* Inspect each component for damage when unpacking. If damage is observed, contact your shipper or service representative immediately.

1. Remove any protective packaging that may be present around the instrument.
2. Examine the packaging material to be sure that the power supply, connecting cables or other components have been removed. The materials that are provided are listed below.
   
   *Note:* Do not discard the packaging material.

### Materials Provided

<table>
<thead>
<tr>
<th>Article</th>
<th>Quantity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hemochron Signature Elite Microcoagulation Instrument</td>
<td>1</td>
</tr>
<tr>
<td>PC Cable (Part No. HJ7405; see <em>Connecting a Computer or Printer</em> on page 24)</td>
<td>1</td>
</tr>
<tr>
<td>AC/DC Power Module (Part No. HX1025)</td>
<td>1</td>
</tr>
<tr>
<td>Operator’s Manual</td>
<td>1</td>
</tr>
<tr>
<td>Hemochron Configuration Manager CD V3.0 or higher</td>
<td>1</td>
</tr>
</tbody>
</table>

*Note:* An AC power cord is provided in the United States, Canada, and Japan only. For other countries, the customer must obtain a power supply cord.

### Materials Required, But Not Provided

<table>
<thead>
<tr>
<th>Article</th>
<th>Quantity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hemochron . Test Cuvettes</td>
<td>As Needed</td>
</tr>
<tr>
<td>Hemochron . Quality Control Products</td>
<td>As Needed</td>
</tr>
</tbody>
</table>

### Optional Materials

<table>
<thead>
<tr>
<th>Article</th>
<th>Quantity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hemochron . PKIT (Printer Kit)</td>
<td>1</td>
</tr>
<tr>
<td>Hemochron . PCKIT (Personal Computer Interface Cable)</td>
<td>1</td>
</tr>
<tr>
<td>Hemochron . CBL-SIG (Serial Printer Interface Cable)</td>
<td>1</td>
</tr>
<tr>
<td>Hemochron <em>ReportMaker™</em> V6.0 or higher CD (Data Management Software)</td>
<td>1</td>
</tr>
<tr>
<td>idms V7.2 or higher (Integrated Data Management System)</td>
<td>1</td>
</tr>
</tbody>
</table>

### BATTERY PRECAUTIONS

Do not open the analyzer. The designed Lithium Ion (Li-Ion) battery is not user replaceable. Opening the analyzer violates the warranty and/or maintenance agreement.

The Hemochron Signature Elite instrument is designed to safely function with the Accriva Diagnostics, Inc. provided Li-Ion battery pack.

Only authorized Accriva service centers are approved to replace the battery pack.

The battery pack is to be charged only with the power supply provided for use with the Hemochron Signature Elite instrument. Refer to section *Charging the Battery* for instructions.

The battery pack may become warm during charging. This is normal.

For optimizing battery life refer to section *Battery Care* for instructions.
In the event of a battery pack leak, do not allow the liquid to come in contact with the skin or eyes. If contact has been made, wash the affected area with water and seek medical advice.
In the event that the battery leaks or is not functioning appropriately, please contact Technical Support to arrange for service of the instrument.
If disposal is necessary, follow local regulations for disposal of Li-Ion batteries.

**Charging the Battery**
The battery of the system should be charged before the system is used.

1. Plug the AC/DC Power Module into an electrical service outlet.
2. Connect the AC/DC Power Module cord to the power connector on the side of the instrument.
   The charge indicator on the instrument keypad will illuminate.
3. Allow the battery to charge for at least sixteen hours.
   
   *Note:* The AC/DC Power Module can remain connected all the time.
4. To ensure adequate charge, leave the instrument connected to the charger for a minimum of sixteen hours. This eliminates the risk of the instrument powering down during a test.

**Battery Warnings**
The message CHARGE BATTERY is displayed intermittently when the remaining battery power is insufficient to ensure test completion. The message BATTERY FAULT is displayed when the battery is completely discharged and cannot be used to run additional tests. The AC/DC Power Module must be used for additional tests until the battery is recharged.
If either the battery voltage or the AC/DC Power Module voltage is too high, the instrument beeps continuously and displays the message DISCONNECT AC ADAPTER IMMEDIATELY. The AC/DC Power Module must be disconnected to shut down the instrument in this condition.

**Checking the Battery**

1. Press and hold **START** to turn on the instrument.
2. Press 0 (zero) to display the main menu:

```
1-Power Status
2-System Time
3-System Date
4-Shut Down
5-System Info
6-Configuration
```
3. Press 1. The battery status is displayed:

   *Note:* Battery status is displayed as **OK**, **LOW**, or **BAD**.
4. Press **CANCEL** twice to return to normal operation.

**Battery Charge Indicators**
A battery charge LED on the front panel (page 5) is illuminated whenever the battery is being charged. The LED flashes when the remaining battery power is low. A battery charge status indicator is also displayed in the upper right corner of the screen when a system disconnected from the AC/DC Power Module is ready to run a test.
Setting the System Time and Date

The time and date a test is run is saved with the test results. The system time and date can be reset by the operator unless the system is in a lockout condition or the time and date are permanently locked (see the Configuration Manager section).

*Note:* The date and time can also be set from the Supervisor menu (page 17) or from Hemochron Configuration Manager (page 69). If the unit has been off for an extended period of time or the battery is very weak, the operator will be prompted to enter or verify that the time and date are correct.

To Set the System Time:
1. Press and hold **START** to turn on the instrument.
2. Press **0** (zero) to display the main menu.
3. Press **2** to display the time:

4. Enter the new time using the numeric keys.

   *Note:* Use the 24-hour format (i.e., enter 15:30 for 3:30 pm). The leading zero is required for single digit times (i.e., enter 06:30 for 6:30 am).
5. Press and hold down **ENTER**. **Stored** is displayed and the entry is saved.
6. Press **CANCEL** to return to normal operation.

To Set the System Date:
1. Press and hold **START** to turn on the instrument.
2. Press **0** (zero) to display the main menu.
3. Press **3** to display the date:

4. Type the new date using the numeric keys.

   *Note:* Use the US (i.e., **MM/DD/YY**) or the IVDD (i.e., **YY-MM-DD**) date format. The leading zero is required for single digit dates (i.e., enter 06/01/06 for June 1, 2006).
5. Press and hold down **ENTER**. **Stored** is displayed and the entry is saved.
6. Press **CANCEL** to return to normal operation.

Instrument Shutdown
1. Press **0** (zero) to display the main menu:

2. Press **4**. The instrument is shut down.

   *Note:* The instrument can also be shut down by pressing and holding down the **START** key for four seconds. The instrument will automatically shut down after being idle for five minutes if there is no cuvette inserted in the test chamber and the instrument is running on the battery.
Displaying System Information
The instrument software version, instrument serial number, and other system software information for the Hemochron Signature Elite instrument can be displayed.

1. Press and hold START to turn on the instrument.
2. Press 0 (zero) to display the main menu:

   ![Main Menu Image](example)

3. Press 5 to display the system information:

   ![System Information Image](example only)

   *Note: The Main Menu will again be displayed after several seconds, or press CANCEL to return to normal operation.*

Configuring Display Brightness, Contrast, and Illumination
The brightness and contrast of the display can be adjusted and the length of time that the display remains illuminated after a test is completed or a key is pressed can be set.

1. Press and hold START to turn on the instrument.
2. Press 0 (zero) to display the main menu.
3. Press 6 for the Configuration menu:

   ![Configuration Menu Image](example)

   **To Adjust the Display Contrast:**
   Display contrast can be set so characters are darker (more contrast) or lighter (less contrast).
   1. Press 1 to display the Contrast menu:
   2. Press 7 to decrease the contrast. Press 9 to increase the contrast.
      *Note: Pressing and holding a key for more than one second will cause it to auto-repeat.*
   3. Press ENTER to save the new setting.

   **To Adjust the Display Brightness:**
   Display brightness can be set so the display background is darker or lighter.
   1. Press 2 to display the Brightness menu:
   2. Press 7 to decrease the brightness. Press 9 to increase the brightness.
      *Note: Pressing and holding a key for more than one second will cause it to auto-repeat.*
   3. Press ENTER to save the new setting.
To Adjust the Length of Time the Display Remains Illuminated:
1. Press 3 to display the Flashlight menu:

```
Flashlight
5 Minutes
ENTER to Save
CANCEL to Exit
```
2. Press 7 to decrease the length of time the display remains illuminated. Press 9 to increase the time.
   
   **Note**: Pressing and holding a key for more than one second will cause it to auto-repeat.

   **Note**: The factory default time for display illumination is five minutes. The maximum illumination time is 100 minutes. To disable display illumination, set the time to zero (0).
3. Press ENTER to save the new setting.

Displaying Available Languages

English is the only language that is available in Hemochron Signature Elite software. Additional languages will be available in future software versions.

To Display the Menu of Available Languages:
1. Press and hold START to turn on the instrument.
2. Press 0 (zero) to display the main menu.
3. Press 6 for the Configuration menu:

```
Configuration
1-Contrast
2-Brightness
3-Flashlight
4-Language Menu
5-Supervisor
```
4. Press 4 to display the menu of available languages:

```
Language Menu
1-English
```
5. Press the numeral key that corresponds to the language to be used.
6. Press ENTER to save the new setting.

Supervisor Functions

The Supervisor menu allows additional actions to be carried out by the laboratory supervisor:
- Scan cuvette or quality control material barcodes to upload the lot number, expiration date, and expected range (for control materials) to Hemochron Configuration Manager
- Scan Operator IDs for uploading to Hemochron Configuration Manager
- Set the instrument date and time (this can also be done from the main menu)
- Print listings of cuvette lot numbers, quality control lot numbers, or Operator IDs
- Set the instrument baud rate
- Enter communication parameters

To Access the Supervisor Functions:
1. Press and hold START to turn on the instrument.
2. Press 0 (zero) to display the main menu.
3. Press 6 for the Configuration menu:

```
Configuration
1-Contrast
2-Brightness
3-Flashlight
4-Language Menu
5-Supervisor
```
4. Press 5. A prompt is displayed to enter the Supervisor PIN:
Note: The Supervisor PIN is set using Hemochron Configuration Manager (see page 53). If an instrument does not yet have a Supervisor PIN, press and hold the ENTER key at the prompt.

5. Enter the Supervisor PIN, then press and hold the ENTER key. The Supervisor menu is displayed:

Scanning Lot Information for Cuvettes or Quality Control Materials
1. On the Supervisor menu, press 1 to display the Scan Lot screen:

2. Hold the instrument over the barcode on the cuvette wrapper or QC package:

Note: Position the scanner port of the instrument approximately four inches from the barcode (see page 5).

3. Press PRINT/SCAN to scan the barcode label. The scanned lot number and a Stored confirmation message are displayed, and the entry is saved.

Note: Additional lots can be scanned by repeating Steps 2 and 3.

4. Press CANCEL to exit.

Scanning Operator IDs
1. On the Supervisor menu, press 2 to display the Scan OID screen:
2. Position the scanner port of the instrument approximately four inches from the barcode (see page 5).
3. Press PRINT/SCAN to scan the barcode. The scanned Operator ID and a Stored confirmation message are displayed, and the entry is saved.
   
   **Note**: Additional Operator IDs can be scanned by repeating Steps 2 and 3.
4. Press CANCEL to exit.

**Setting the System Time or System Date**

For convenience, the Supervisor menu includes the same commands that are on the main menu for changing the system time and system date.

1. On the Supervisor menu, press 3 to set the system time.
2. On the Supervisor menu, press 4 to set the system date.
   
   **Note**: Procedures for setting the system time and system date are described on page 15.

**Specifying the Time Zone**

The time zone setting is used to compare the instrument time to Greenwich Mean Time (GMT). This allows the instrument to communicate with other devices in accordance with CLSI Standard POCT-1A.

1. On the Supervisor menu, press 5 to display the Time Zone screen:

   ![Time Zone Screen]

2. Enter the difference (in hours and minutes) between the local time zone and GMT. For example, if the local time is 2:00 pm and GMT is 7:00 pm, enter -5.00.
   
   **Note**: Press the 8 key to enter positive (+) time differences in 15-minute increments. Press the 0 key to enter negative (-) time differences in 15-minute increments.
3. Press ENTER to store the time zone.
4. Press CANCEL to exit.

**Specifying Print Options**

Listings of cuvette lot numbers, quality control lot numbers, or Operator IDs can be printed at an external printer (see page 24 for instructions to connect the printer).

1. On the Supervisor menu, press 6 to display the Print Options menu:

   ![Print Options Menu]

2. Press 1 to print a listing of Operator IDs.
3. Press 2 to print a listing of cuvette lot numbers.
4. Press 3 to print a listing of quality control material numbers.

**Specifying the Baud Rate**

Listings of cuvette lot numbers, quality control lot numbers, or Operator IDs can be printed at an external printer (see page 24 for instruction to connect the printer).

1. On the Supervisor menu, press 6 to display the Print Options menu:
2. Press 4 to display the Baud Rate screen.

3. Press the numeral key that corresponds to the baud rate to be selected.

   Note: The baud rate that is currently specified is designated by an asterisk (*).

4. Press CANCEL to exit.

Specifying the Communication Protocol

The instrument can communicate with a personal computer (using the COM port) or a network (using the Ethernet port). One of two communication protocols can be selected:

Accriva Diagnostics, Inc. proprietary communication protocol that allows records to be transferred from the instrument and allows the instrument to be configured using Accriva Diagnostics, Inc. software such as Hemochron Configuration Manager. This protocol is supported over COM or NET.

A communication protocol that meets CLSI POCT-1A standards. This protocol allows the instrument to communicate with any software or device that is compliant with POCT-1A. This protocol is supported over NET only. If Port COM is selected, the following message is displayed:

```
UNSUPPORTED
POCT >> COM
```

Note: If the CLSI POCT-1A compliant communication protocol is used, communication must be initiated from the instrument. If the Accriva proprietary communication protocol is used, communication must be initiated from either the personal computer or the network.

1. From the Supervisor menu, press 7 to display the Communications menu:

2. Press 1 to select the desired communication protocol. (Display OFF to use the Accriva proprietary communication protocol, or display ON to use the CLSI POCT-1A compliant communication protocol.)

   Note: If Port is set to COM when attempting to set POCT ON, the setting will not be allowed and the message UNSUPPORTED POCT >> COM will be displayed. Specifying POCT connectivity acts as a toggle. If CLSI POCT-1A connectivity standards are already specified (POCT ON is displayed), this can be cancelled by pressing 1 again to display POCT OFF.

Specifying the Communication Port

1. From the Supervisor menu, press 7 to display the Communications menu:
2. Press 2 to select the desired port. (Display COM to specify use of the COM port, or display NET to specify use of the Ethernet port.)

*Note:* If POCT is set to ON when attempting to set Port COM, the setting will not be allowed and the message UNSUPPORTED POCT >> COM will be displayed. Specifying the port acts as a toggle. If use of the Ethernet port is already specified (Port NET is displayed), this can be cancelled by pressing 2 again to display Port COM.

**Specifying Continuous POCT-1A Communication**

*Note:* Communication is initiated from the instrument only when the CLSI POCT-1A compliant communication protocol is used. If the Accriva proprietary communication protocol is used, communication is initiated from a personal computer or network.

1. From the Supervisor menu, press 7 to display the Communications menu:

2. Press 3 to select the desired communication option when the CLSI POCT-1A compliant communication protocol is used. (Display OFF to initiate communication only if requested, or display ON to initiate continuous communication whenever the instrument is started.)

*Note:* Specifying the communication option acts as a toggle. If initiation of communication upon startup is already specified (Push POCT ON is displayed), this can be cancelled by pressing 3 again to display Push POCT OFF.

**Entering Network Communication Parameters**

Some or all of the following parameters may need to be entered in order for the instrument to communicate over a network:

- IP Address
- Gateway
- NetMask
- Remote IP
- Remote Port

*Note:* Consult with the network administrator for your facility to determine which parameters must be entered and what those settings should be.

1. From the Supervisor menu, press 7 to display the Communications menu:

2. (If the CLSI POCT-1A compliant communication protocol is currently selected): Press 1 to toggle the communication protocol to the Accriva proprietary communication protocol:
Note: The CLSI POCT-1A compliant communication protocol must be OFF when running XPORT Setup.

3. Press 4 to start the XPORT Setup module. The message XPORT Setup Please Wait is displayed, followed by the first page of the XPORT Setup module:

![First Page](image)

Note: If the message XPORT Setup NOT RESPONDING! is displayed, contact Technical Support at (877) 579-4441 (toll free) or (858) 263-2502, or by e-mail at techsupport@accriva.com.

4. Press 0 to display the next page (for entering IP Address and Gateway settings):

![Next Page](image)

5. Press 1 to display the screen for entry of the IP Address:

![IP Address](image)

Enter the IP Address using the numeral keys:

![IP Address](image)

Note: Press Enter to move to the next field without entering a numeral. Press Cancel to move to the previous field.

6. After the IP Address has been entered, press and hold ENTER. The message Parameter String Stored is displayed, followed by the screen for entering IP Address and Gateway settings:

![Parameter String Stored](image)

Important: The parameter string is temporarily stored. Even though the new parameter is displayed on the screen, it is not saved until Enter is pressed to exit XPORT Setup (Step 13).

Note: A range of IP addresses (from 169.254.0.1 to 169.254.255.1) has been reserved for Auto-IP enabled devices. This range of Auto IP addresses should not to be used over the Internet.

Auto-IP can be disabled by setting the device’s IP address to 000.000.001.000. This setting enables DHCP but disables Auto-IP.

7. Press 2 to display the screen for entry of the Gateway. Enter the Gateway in the same manner as for the IP Address, then press and hold ENTER to again display the screen for entering IP Address and Gateway settings.

8. Press 0 to display the next page (for entering NetMask and Remote IP settings) of the XPORT Setup module:
9. Press 3 to display the screen for entry of the NetMask. Enter the NetMask in the same manner as for the IP Address, then press and hold ENTER to again display the screen for entering NetMask and Remote IP settings.

10. Press 4 to display the screen for entry of the Remote IP. Enter the Remote IP in the same manner as for the IP Address, then press and hold ENTER to again display the screen for entering NetMask and Remote IP settings.

11. Press 0 to display the next page (for entering the Remote Port setting) of the XPORT Setup module:

12. Press 5 to display the screen for entry of the Remote Port. Enter the Remote Port in the same manner as for the IP Address, then press and hold ENTER to again display the screen for entering the Remote Port setting.

13. Press ENTER while any page of the XPORT Setup module is displayed to save the new parameters. The message Updating XPORT is displayed, followed by the Communications menu (see Step 1).

   **Important**: Parameters are temporarily stored until the Enter key is pressed while any page of the XPORT Setup module is displayed. If the Cancel key is pressed while any page of the XPORT Setup module is displayed, the message Cancel without updating XPORT? is displayed for confirmation by the operator.

   **Note**: The parameters can be printed to a device connected to the COM port by pressing the PRINT/SCAN key.

**Configuring Network Communication Parameters Using a Personal Computer**

An experienced network administrator can communicate with the XPORT Setup module and enter communication parameters using a personal computer connected to the COM port.

1. From the Supervisor menu, press 7 to display the Communications menu:

2. (If the CLSI POCT-1A compliant communication protocol is currently ON): Press 1 to toggle ON to OFF for the proprietary communication protocol:

   **Note**: The CLSI POCT-1A compliant communication protocol must be OFF when running XPORT Setup.

3. Press 5 to start the XPORT COM-1 Setup module. The message XPORT Setup Please Wait is displayed, followed by the message XPORT COM1-SET.

   The XPORT module can now communicate with the personal computer through the COM port, using terminal emulator software such as Microsoft Windows Hyperterminal.
4. Upon completing the communication session, press **Cancel** to exit the session and display the Communications menu again.

**Connecting a Computer or Printer**

An IBM-compatible laboratory computer or a serial printer (Seiko DPU-414 is recommended) can be used with the system. The connector port for these devices is located at the side of the instrument.

![Connector Ports](connector-ports.png)

**To Connect to a Printer:**

1. Obtain a printer interface cable.
   
   **Note:** The printer interface cable is included with the optional Hemochron . PKIT (Printer Kit) only. A separate CBL-SIG (Serial Printer Interface Cable) can be obtained from Accriva Diagnostics, Inc., or a cable can be prepared as outlined below. If a printer is connected, the serial printer communication parameters must be set to 9600 baud with no parity, 8 data bits and 1 stopbit using XON/XOFF 3-wire software protocol. For additional information on printer configuration, contact Technical Support at (877) 579-4441 (toll free) or (858) 263-2502, or by e-mail at techsupport@accriva.com.

2. Connect one end of the cable to the printer connector port on the side of the instrument.

3. Connect the other end of the printer cable to the printer.

**To Connect to a Personal Computer:**

1. Obtain a computer interface cable.
   
   **Note:** The computer interface cable is included with the instrument. A replacement PCKIT (Personal Computer Interface Cable) can be obtained from Accriva Diagnostics, Inc., or a cable can be prepared as outlined below. Contact Technical Support for instructions on setting up the personal computer.

2. Connect one end of the cable to the computer connector port on the side of the instrument.

3. Connect the other end of the cable to an unused serial communication port on the computer.

**To Connect to a Network:**

1. Obtain a network interface cable.

2. Connect one end of the cable to the Ethernet port on the side of the instrument.

3. Connect the other end of the cable to a network connection port.
Preparing a Printer or Computer Interface Cable

Cabling and connectors can be purchased locally at an electronics store. Use an 8-wire RJ45 to RJ45 modular straight through connecting cable (no longer than 25 feet) plus an RJ45 to DB9 adapter. Numbering of the pins on the RJ45 connectors is shown below:

The cable configurations used for connecting a serial printer or a computer are shown below:

Connecting the instrument to the Seiko DPU-414 printer with a DB9S connector

Connecting the instrument to any personal computer with a DB9P connector

STARTING THE INSTRUMENT

Start the instrument by pressing and holding down the START key or by inserting a cuvette:

Note: Allow cuvettes to reach room temperature (15 to 30 °C) before opening the pouch. This may take up to 60 minutes. Consult the cuvette package insert for additional information concerning cuvette storage and handling.

Note: Insert the cuvette with the label facing up. Push the cuvette all the way in until it stops.

Starting the Instrument by Pressing the START Key

1. When the START key is pressed, the following screens are briefly displayed:
2. The instrument then prompts the user that the instrument is ready to accept a cuvette:

Starting the Instrument by Inserting a Cuvette
1. When a test cuvette (e.g., ACT+) is inserted to turn on the instrument, the following prompts are briefly displayed:

   
   (example only)

   

   Followed by

   

   Note: The test name for the inserted cuvette is displayed in the upper left-hand corner of the screen. Check the display for fault messages while the cuvette is warming. Refer to the Troubleshooting section for the procedure to resolve a fault message.

2. The instrument then warms the cuvette to 37 °C ±1.0 °C:

3. When the cuvette reaches temperature, the instrument beeps and alternately displays:

4. The sample can now be added. After the sample is added, press START to run the test.

RUNNING A TEST

Samples are designated as patient samples by default. If a control is being run, the sample type must be changed from patient sample to QC Normal or QC Abnormal. If a patient sample is being run, this step is unnecessary.

Important: Failure to select QC Normal or QC Abnormal for a control sample will result in storage of the control results as patient results in the patient database.

Designating the Sample as a Normal or Abnormal Control
1. Press QC to display the QC SELECTS menu:

   

   Note: A cuvette must be inserted into the instrument to display the QC SELECTS menu. If entry of a cuvette lot number and/or Operator ID or PIN is required, this must be done before the QC SELECTS menu can be displayed.

2. Press 1 to designate the sample as a normal control (QC Normal is displayed), or press 2 to designate the sample as an abnormal control (QC Abnormal is displayed).
Note: The sample type can be designated either before or during the test, unless QC lockout features are enabled. Once QC is selected, pressing the QC key again briefly displays the QC level and assay being run.

To exit the QC menu, press the CANCEL key.

Important: Once a QC type is selected, it cannot be changed.

Entering an Optional Patient ID (PID) and/or an Operator ID (OID)
A Patient ID (PID) and/or an Operator ID (OID) can be entered for each test. The IDs will be stored with the test results. Entry of a PID and/or OID is optional unless a mandatory entry is specified (see the Configuration Manager section on page 54).

The PID and OID can either be read from a barcode label using the internal barcode scanner or can be entered using the keypad (unless entry of PID and OID by barcode scanning is specified in Configuration Manager). Any leading or trailing spaces will be ignored when scanned.

Note: Any PID or OID that is manually entered or accepted for Reuse will be flagged as a manual entry in the database. Backspace and overwrite characters as needed, or backspace to the first entry and then backspace twice to clear all entries.

To Enter a Patient ID:
1. Select ID to display the ID SELECTS menu:

Note: A cuvette must be inserted into the instrument to display the ID SELECTS menu.

2. Press 1 to display the screen for entering the PID:

Note: An acceptable PID is any combination of 1 to 20 characters. During PID entry, a PID exceeding 16 characters will wrap to the next line. The number of characters to be entered for PID (from 1 to 20) may be designated via Configuration Manager. An acceptable range of PID lengths may also be designated. Check the PID by pressing ID and then pressing 1 to display the entered value.

3. Scan the PID from a barcode label using the internal barcode scanner, or enter it using the keypad.
   a. To scan the PID from a barcode label, position the scanner port approximately four inches from the barcode label (see page 5) and then press PRINT/SCAN. The scanned PID and a Stored confirmation message are displayed:

   b. To enter the PID manually, type the PID using the keypad and then press and hold ENTER. The entered PID and a Stored confirmation message are briefly displayed as shown above.

   Note: Numeric and/or alphabetic characters can be entered. Press the NOTE key to change from numeric to alphabetic characters or vice versa.
If alphabetic characters are being entered, advance from the displayed letter to the next letter on that key by pressing the key again before the cursor advances.

Note: The PID can be designated (or changed) anytime. Check the PID by pressing ID and then pressing 1 to display the entered value.

To Enter an Operator ID:

1. Select ID to display the ID SELECTS menu:

Note: A cuvette must be inserted into the instrument to display the ID SELECTS menu.

2. Press 2 to display the screen for entering the OID:

3. Scan the OID from a barcode label using the internal barcode scanner, or enter it using the keypad.
   a. To scan the OID from a barcode label, position the scanner port approximately four inches from the barcode label (see page 5) and then press PRINT/SCAN. The scanned OID and a Stored confirmation message are displayed:
      Note: For software versions 2.2 and higher, the Elite uses a truncating feature to scan an operator ID greater than 16 characters. When scanning an operator ID greater than 16 characters, the first 16 characters will be accepted. Any additional characters beyond the first 16 will not be stored. For software versions below 2.2, when scanning operator IDs greater than 16 characters, an error message will be displayed and ID may be entered manually as stated below.
      b. To enter the OID manually, type the OID using the keypad and then press and hold ENTER. The entered OID and a Stored confirmation message are displayed as shown above.
      Note: Up to 16 numeric and/or alphabetic characters can be entered. Press the NOTE key to change from numeric to alphabetical characters or vice versa. If alphabetic characters are being entered, advance from the displayed letter to the next letter on that key by pressing the key again before the cursor advances.
      Note: The OID can be designated (or changed) anytime unless operator lockout is enabled through Configuration Manager. Check the OID by pressing ID and then pressing 2 to display the entered value.

Entering a Mandatory Cuvette Lot Number and Expiration Date

Entry of the cuvette lot number and expiration date can be required for completion of a test.

Note: Prompts for entry of cuvette lot information are only displayed if the appropriate cuvette lot option is enabled through Configuration Manager (see page 54). If Required, Scan Only, and/or Pre-stored Only are selected as a cuvette lot option, the corresponding prompts to enter the information are automatically displayed. If none of the cuvette lot options are selected, there is no other means for cuvette lot information to be entered.

To Enter Mandatory Cuvette Information:

1. The prompt to enter a cuvette lot number is automatically displayed when a cuvette is inserted:
2. Select the cuvette lot number from the list, scan the cuvette lot number, or enter it using the keypad.
   a. To select the cuvette lot number from the list, press the numeral key for that lot number. For example, press 2 to select lot number L5NCS055 in the example above.
      Note: Information for up to five lots of cuvettes for each assay type can be scanned using the Supervisor menu (page 17) or entered using Configuration Manager (see page 61).
      A cuvette lot number that has been scanned or entered is not displayed on the list after the lot reaches its expiration date. If more than five cuvette lot numbers per assay are scanned or entered, the lot number(s) with the closest expiration date will be deleted.
   b. To scan the cuvette lot number and information, **position the scanner port approximately four inches from the barcode** (page 5) on the cuvette wrapper and then press PRINT/SCAN.

   ![Barcode and Cuvette Information]

   The scanned lot number and **Stored** are displayed.

   c. To enter the cuvette lot number and information manually, press 7 to display the screen for entering the cuvette lot number:

   ![Enter Cuv Lot]

   Enter the cuvette lot number using the numeral keys:

   ![Enter Cuv Lot]

   **Note:** Press the NOTE key to change from numeric to alphabetical characters or vice versa. When alphabetic characters are being entered, advance from the displayed letter to the next letter on that key by pressing the key again before the cursor advances.
   **Note:** Cuvette lot numbers that are manually entered into the Hemochron Signature Elite instrument are only saved to the current test record. They are not uploaded to Configuration Manager.

   After the cuvette lot number has been entered, press and hold **ENTER** to display the screen for entering the cuvette information:

   ![Cuv Info Field]

   Enter the cuvette information using the numeral keys:
Entering a Mandatory PID and/or OID or PIN

Entry of a Patient ID (PID) and/or an Operator ID (OID) or Operator Personal Identification Number (PIN) can be required for completion of a test (see the Configuration Manager section on page 54). If entry of one or more of these IDs is required, appropriate prompts for entry of PID and/or OID or PIN are automatically displayed.

To Enter a Mandatory Patient ID:

1. The PID prompt is automatically displayed when a cuvette is inserted (or after a mandatory cuvette lot and/or OID is entered):

   ![PID prompt example](image)

   Note: The dashed line after the cursor denotes the minimum length of the PID. The dotted line following the dashed line denotes the maximum length of the PID (if specified in Configuration Manager).

   Note: Pressing the QC button after the PID prompt is displayed allows a QC test to be performed. Entry of a PID is not required in this case.

2. Scan the PID from a barcode label using the internal barcode scanner, or enter it using the keypad.

   Note: Refer to Step 3 on page 27 for the procedure to scan the PID using the barcode scanner or to enter it manually.

3. Press and hold ENTER. If an acceptable PID is entered, operation of the instrument continues:

   ![PID entered successfully](image)

   Note: When running a test or viewing results, a PID exceeding 11 characters will wrap to the next line.

To Enter a Mandatory Operator ID:

1. The OID prompt is automatically displayed when a cuvette is inserted (or after a mandatory cuvette lot and/or QC lot is entered):

   ![OID prompt example](image)

2. Scan the OID from a barcode label using the internal barcode scanner, or enter it using the keypad.

   Note: Refer to Step 3 on page 28 for the procedure to scan the OID using the barcode scanner or to enter it manually.

3. Press and hold ENTER. If an acceptable OID is entered, operation of the instrument continues:

   ![OID entered successfully](image)
Note: If entry of any OID is specified, an acceptable OID is any number up to 16 digits. If entry of a valid OID is specified, an acceptable OID is any number contained in the Operator Table (see page 59).
Check the OID by pressing ID and then pressing 2 to display the entered value.

4. If an unacceptable OID is entered, ID is Not Valid is displayed followed by the OID prompt. Operation of the instrument will not continue until an acceptable OID is entered.

To Enter a Mandatory Operator PIN:
1. The PIN prompt is automatically displayed when a cuvette is inserted:

2. Enter the PIN using the keypad.
3. Press and hold ENTER. If an acceptable PIN is entered, operation of the instrument continues:

   followed by

   Note: An acceptable PIN is any string of up to nine numeric digits (other than 0) that matches a PIN in the Operator Table.

4. If an unacceptable PIN is entered, ID is Not Valid is displayed followed by the PIN prompt. Operation of the instrument will not continue until an acceptable PIN is entered.

Prompts for Mandatory QC Tests
Electronic quality control (EQC) tests and/or liquid quality control (LQC) tests can also be required to be run at specified intervals (see the Configuration Manager section). When it is time for a required QC test to be run, the required QC test must be successfully completed before the specific patient test(s) can be run. If EQC is required, no patient tests can be run until successful completion of EQC.

Note: A specified limited number of additional tests per assay (911 exceptions) can be run after a mandatory QC interval has been reached. Successful QC tests for an assay will automatically reset the 911 exception for that assay. Consult the Configuration Manager section for details.

To Run a Mandatory QC Test:
1. After a specified QC interval (see page 55 in the Configuration Manager section) passes without a successful QC test, a prompt to run the required QC test is automatically displayed:

2. Press the QC key to display additional information concerning QC testing that is required (REQ):

3. If a cuvette is inserted for a test with an expired QC interval, the message LQC Expired is displayed, followed by a prompt to run either a QC test or another patient sample (if the number of 911 Exceptions has not been exceeded):
4. Run the QC test according to the displayed prompts. If the results for the QC test are acceptable, patient tests can now be run.

**Entering a Mandatory Lot Number, Expiration Date, and Expected Range for Controls**

Entry of the lot number, expiration date, and expected range of liquid controls can be required for completion of a QC test.

*Note:* Prompts for entry of the lot number, expiration date, and expected range of liquid controls are only displayed if the appropriate QC lot option is enabled through Configuration Manager (see page 55). If Required, Scan Only, and/or Pre-stored Only are selected as a QC lot option, the corresponding prompts to enter the information are automatically displayed. If none of the QC lot options are selected, there is no other means for QC lot information to be entered.

**To Enter Mandatory Control Information:**

1. The prompt to enter a QC lot number is automatically displayed when a QC test is selected:

2. Select the QC lot number from the list, scan the control lot number, or enter it using the keypad.
   a. To select the QC lot number from the list, press the numeral key for that lot number. For example, press 1 to select lot number E5DCA007 in the example above.
      *Note:* Information for each lot of controls is entered using Configuration Manager (see page 62). Information for up to five lots of normal controls and five lots of abnormal controls can be stored for each assay type. A control lot number that has been scanned or entered is not displayed on the list after the lot reaches its expiration date.
   b. To scan the control information, **position the scanner port approximately four inches from the barcode** (see page 5) on the liquid control packaging and then press PRINT/SCAN.
   c. To enter the QC lot number and information manually, press 7 to display the screen for entering the QC lot number:

Enter the QC lot number using the numeral keys:
**Note:** Press the NOTE key to change from numeric to alphabetical characters or vice versa. When alphabetic characters are being entered, advance from the displayed letter to the next letter on that key by pressing the key again before the cursor advances.

**Note:** QC lot numbers that are manually entered into the Hemochron Signature Elite instrument are only saved to the current test record. They are not uploaded to Configuration Manager.

After the QC lot number has been entered, press and hold ENTER to display the screen for entering the QC information:

Enter the QC information using the numeral keys:

**Note:** The QC information field consists of numerical characters (1406122890515200 in the example). The expiration date and expected range of the control is encoded in the QC information.

After the QC information has been entered, press and hold ENTER to continue with the test.

### Adding a Note

Up to two operator-selectable notes can be added to the record for each test and will be printed with the final record. The notes are selected from a menu of nine available notes.

**Note:** Consult the Configuration Manager section for the procedure to add new user-selectable notes to the Note Table.

1. Press NOTE:

   ![Note](image)

   **Note:** A cuvette must be inserted into the instrument before a note can be entered.

2. Type a number (1 through 9) to display the corresponding note in the first note field. For example:

   ![Note](image)

3. To enter a note in the second note field, press ENTER to highlight the second note field before typing a number for the second note:
Note: No Note is displayed if a note is not defined for a note field. To remove a currently entered note, highlight the note field and enter 0 to display No Note.

Note: If a third note is entered, the first note is cancelled.

4. Press CANCEL. The selected note (or notes) are entered into the test record.

Specimen Collection

Collect blood specimens according to CLSI document H21-A5, entitled Collection, Transport and Processing of Blood Specimens for Coagulation Testing and General Performance of Coagulation Assays.

Important: Collect blood specimens in a manner that prevents contamination with tissue thromboplastin, indwelling intravenous (I.V.) solutions, or alcohol cleansing solution. Discard samples that are not properly collected or contain visible clots or debris. Use a 23 gauge or larger needle if a syringe is used for blood collection. If a sample is expelled through the same needle, do it slowly to prevent hemolysis. Consult the cuvette package insert for additional information concerning specimen collection and storage.

Starting the Test

After initialization is completed the instrument beeps once, signaling the instrument is ready for the sample. Add Sample and Press Start are displayed with a countdown to START TIMED OUT (see the note below), and the blood sample can be placed into the cuvette and the test started.

Note: START TIMED OUT is displayed if the test is not started within five minutes. If this occurs, repeat the test using a new cuvette.

1. Place a drop of blood into the sample well of the cuvette:

   Note: Fill the sample well from the bottom up. Add enough blood so that the lower wall of the center sample well is completely filled. If the meniscus of the blood sample extends above the wall, push the excess blood into the overflow area. A transfer needle can be used to transfer blood.

2. Press the START key. The elapsed time (in seconds) is displayed until the sample clots.

   Note: SAMPLE TOO SMALL, SAMPLE TOO LARGE, or SAMPLE NOT SEEN is displayed if an incorrect amount of sample was added. Repeat the test using a new cuvette.

3. When a clot is detected, the instrument beeps once. Final results of the test are calculated and displayed.

   Note: The whole blood clotting times are converted to plasma equivalent values (for APTT and APTT Citrate), or plasma equivalent values and International Normalized Ratio (for PT and PT Citrate). Celite equivalent values are displayed for ACT+ and ACT-LR.

4. Remove the cuvette and discard it.

   Note: Results are saved in the database and printed if a printer is connected. The date and time of the test, cuvette lot and/or QC lot information, PID, OID, note(s) (if entered), and error message (if a fault is detected for the test or sample) are printed with the test result.
QUALITY CONTROL (QC)

The Joint Commission on Accreditation of Healthcare Organizations (JCAHO) and other regulatory agencies recommend that medical and laboratory instrumentation be enrolled in a quality assurance program adequate in maintaining accurate and reliable performance of the equipment. Complete records of such quality control must be kept.
Routine quality control testing should be part of a comprehensive quality assurance program.
Quality control testing of the Hemochron Signature Elite consists of the following operations:

- Testing system performance and verifying system temperature using internal Electronic System Verification every eight hours of operation.
- Testing cuvettes in accordance with the Package Insert for each assay using two levels of liquid controls.

Self-Check

The Hemochron Signature Elite instrument performs a “self-check” every time it is activated and a test is performed. When a test is initiated by inserting a cuvette, system checks are automatically performed and include:

- Verification of adequate battery power to complete a full test.
- Verification of the test type on the screen display to insure that the LEDs used for identifying the test are functioning properly.
- Verification that the cuvette temperature is warmed to 37 °C ±1.0 °C. If this temperature is not achieved or is exceeded, an appropriate error message will be displayed and testing is prohibited.
- Verification that the sample is present and is of sufficient size to run test. This insures that the pumps and sample sensing LEDs are functioning properly and that the cuvette is adequately sealed. If these instrument and sample parameters are not appropriate, the test is terminated and an error message displayed.
- Verification that the internal timers function correctly for each test. If the system timer and assay timer disagree, a real-time clock error message is displayed and the test result is not reported.

Verification of Instrument Temperature

Instrument temperature is automatically checked whenever EQC is carried out. The instrument temperature check verifies that a temperature of 37 °C ±1.0 °C is maintained during the test.

**Note:** If the temperature reading is out of range, please contact Technical Support at (877) 579-4441 (toll free) or (858) 263-2502.

QC of Instrument Performance

The instrument should be tested at two levels once every eight hours of operation. Automatic internal Electronic Quality Control (EQC) can be used to provide a two-level electronic verification of instrument performance, or liquid quality control products can be used.

**Note:** Hemochron Configuration Manager is used to specify Automatic EQC, to specify the required interval for performing EQC, and to program EQC Level 2 for either 300 seconds or 500 seconds (see page 58).

**Important:** If Auto EQC is selected, ensure that cuvettes are routinely removed from the instrument after completing a test. If EQC is run while a cuvette is inserted, the message Detector Blocked is displayed and the EQC test fails.

EQC is performed automatically if Automatic EQC is specified and if the Hemochron Signature Elite instrument is plugged into the AC/DC Power Module. EQC results are stored to the database and displayed for five minutes (or until Cancel is pressed).

**QC Using EQC:**

1. Display the QC Status menu:
Note: Display the QC Status menu by pressing the QC key before a cuvette is inserted. If EQC is run while a cuvette is inserted, the message Detector Blocked is displayed and the EQC test fails.

2. Press 1. The test chamber warms to temperature and the EQC test begins. The results are displayed while the test is progressing.

3. When the test is completed, the results are displayed on the screen and written to the QC database. Press CANCEL to exit the screen.

Note: Internal EQC will check two levels of QC and the temperature, and it will store each result. If one result fails, the test will stop and record all results as failed. If the user aborts the internal EQC test, the test is not saved to the database or printed.

QC Using Liquid Quality Control Products:
1. Obtain the liquid quality control products.
   
   Note: Hemochron . Quality Controls are recommended. Refer to the package insert for the liquid quality control products for storage and handling information.

2. Obtain a cuvette for the test to be validated and insert it into the instrument.

3. To run QC when it is not required, press the QC key.

4. Select QC NORMAL (or QC ABNORMAL) and enter an Operator ID (optional) for the test.

   Important: LQC results will be stored in the patient database if QC NORMAL or QC ABNORMAL is not selected for the test.

5. When Add Sample and Press Start are displayed, add the appropriate liquid control sample and run the test in the same manner as for a patient sample.

   Note: If any error messages appear on the instrument screen, remove the cuvette and repeat the test using a new cuvette.

QC of Cuvettes

Cuvette validation is carried out using the appropriate Hemochron Quality Control product and the test procedure provided.

Acceptable performance ranges for the Hemochron . cuvettes are included with each Hemochron QC product. The cause of out-of-range results is likely attributable to test technique, control material, cuvette and/or the instrument. Refer to the Troubleshooting section for additional information.

Checking QC Intervals

If EQC tests and/or LQC tests are required to be run at specified intervals, the amount of time remaining before QC must be run on each test can be checked.

To Check QC Intervals:

   Note: If a cuvette is not inserted in the instrument, pressing QC will immediately display QC Status and will allow the operator to run EQC.

1. With a cuvette inserted, press QC to display the QC SELECTS menu:
2. Press 4. The remaining time (shown as **hours:minutes**) before QC must be run on each test is displayed. Tests are listed in the order which QC will need to be run, with EQC always displayed last:

<table>
<thead>
<tr>
<th>QC Status</th>
<th>Press 1-Run EQC</th>
</tr>
</thead>
<tbody>
<tr>
<td>PT cit</td>
<td>10:00</td>
</tr>
<tr>
<td>ACT-LR</td>
<td>13:20</td>
</tr>
<tr>
<td>APTT</td>
<td>18:20</td>
</tr>
<tr>
<td>aPTT</td>
<td>20:30</td>
</tr>
<tr>
<td>PT</td>
<td>30:30</td>
</tr>
</tbody>
</table>

3. If applicable, the screen will toggle to display the QC time remaining for additional tests.

<table>
<thead>
<tr>
<th>QC Status</th>
<th>Press 1-Run EQC</th>
</tr>
</thead>
<tbody>
<tr>
<td>ACT+</td>
<td>32:30</td>
</tr>
<tr>
<td>EQC</td>
<td>12:20</td>
</tr>
</tbody>
</table>

**OPERATING PRECAUTIONS**

DO NOT open the instrument, as there are no user-serviceable parts and the warranty is violated.

DO NOT remove the AC/DC Power Module from the instrument by pulling on the cord.

DO NOT use cuvettes that are past their marked expiration date or have been improperly stored. Only use Hemochron Signature Elite cuvettes.

DO NOT force a cuvette into the instrument. If resistance to insertion is encountered, gently remove the cuvette and examine the cuvette slot. Remove any obstruction before attempting further use of the instrument (see *Service and Maintenance* on page 76).

DO NOT use excessive force in pressing the instrument keys.

DO NOT expose the instrument to extreme temperatures (above 50 °C).

DO NOT drop the instrument, and do not use the results if the instrument is dropped during a test. Use of an AC adapter other than that provided with the instrument could lead to reduced safety or instrument damage.

The use of accessory equipment identified in this manual either in the patient vicinity, or that does not comply with either the equivalent safety requirements of this equipment or IEC 61010, may lead to a reduced level of safety in the resulting system.

The Hemochron Signature Elite instrument should only be used by healthcare professionals trained and certified in the use of the system and operated in accordance with facility policies and procedures.

All biohazard safety guidelines pertaining to the handling and disposal of human blood should be strictly adhered to when collecting and handing blood specimens and when operating the Hemochron Signature Elite Microcoagulation instrument.

Used Hemochron Signature Elite test cuvettes should be considered as potentially infectious. They should be handled according to individual institutional policies concerning the disposal of potentially infectious materials.

**CAUTION:** Verify that any third-party connectivity software to be used is compatible with the software version of the Hemochron Signature Elite instrument in use. Transferred data will be lost if third-party connectivity software used is not compatible with Hemochron Signature Elite software.
LIMITATIONS
Test results of the Hemochron Signature Elite Microcoagulation instrument are affected by poor technique during blood collection and delivery to the sample well. The accuracy of the test is largely dependent upon the quality of the blood specimen, which is dependent upon the blood sample collection and the transfer of blood to the test cuvette. Refer to the individual assay package insert for specific limitations. Hemochron Signature Elite test results should always be scrutinized in light of a specific patient’s condition or anticoagulant therapy. Any test results exhibiting inconsistency with the patient’s clinical status should be repeated or supplemented with additional diagnostic tests. Samples with a hematocrit less than 20% or greater than 55% are not recommended, due to an optical density outside the detection level of the Hemochron Signature Elite Microcoagulation instrument.

RESULTS MANAGEMENT
Up to 600 patient test results and 600 quality control test results can be stored in the instrument. In addition to test results, the date and time of each test, PID, OID, and user notes are also stored.

Note: PID and/or OID are not stored if a value is not entered when the test is run. A PID is not allowed for any QC test.

Note: If an error is encountered with the test or the sample, the error message for the error will be included on the printout. Refer to page 72 for error messages, their cause, and their resolution.

Printing Results
1. Press PRINT/SCAN to display the PRINT menu:

   Then perform one of the following procedures:

   **To Print Results of the Last Test:**
   1. Press 1. Results of the last test are printed.

   **To Print All Results in a Database:**
   1. Press 2 to print all of the results in the patient database. Or, press 3 to print all of the results in the QC database.

   **To Print Patient Results:**
   1. Press 4 to display the menu for printing patient results:

      Then perform the desired tests:
      a. To print results for a specific patient, press 1 to display the Print by PID screen:

      Scan or type the PID, then press ENTER. All test results for the specified patient are printed.
b. To print results obtained by a specific operator, press 2 to display the Print by OID screen. Scan or type the OID, then press ENTER. All patient test results obtained by the specified operator are printed.

c. To print results obtained on a specific date, press 3 to display the Print by Date screen:

![Print by Date](image)

Enter the date using the keypad, then press ENTER. All patient test results obtained on the specified date are printed.

d. To print all results for a specific assay, press 4 to display the Print by Assay screen:

![Print by Assay](image)

Note: If the desired assay is not shown, press 7 to display additional assays. Enter the number for the assay, then press ENTER. All patient test results for the specified assay are printed.

e. To print results for a specific cuvette lot number, press 5 to display the Print by CVlot screen:

![Print by CVlot](image)

Scan or type the cuvette lot number, then press ENTER. All patient test results for the specified cuvette lot number are printed.

f. Press CANCEL to return to the menu for printing patient results (Step 1).

To Print QC Results:

1. Press 5 to display the menu for printing QC results:

![BY QC MENU](image)

2. Print the results for the desired tests:

a. To print QC results obtained by a specific operator, press 1 to display the Print by OID screen:

![Print by OID](image)

Scan or type the OID, then press ENTER. All QC test results obtained by the specified operator are printed.
b. To print results obtained on a specific date, press 2 to display the Print by Date screen:

```
<table>
<thead>
<tr>
<th>Print by Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hold ENTER=Store</td>
</tr>
<tr>
<td>Hold CANCEL=Exit</td>
</tr>
</tbody>
</table>
```

Enter the date using the keypad, then press ENTER. All QC results obtained on the specified date are printed.

c. To print all results for a specific assay, press 3 to display the Print by Assay screen:

```
<table>
<thead>
<tr>
<th>Print by Assay</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
</tr>
<tr>
<td>B</td>
</tr>
<tr>
<td>1</td>
</tr>
<tr>
<td>2</td>
</tr>
<tr>
<td>3</td>
</tr>
<tr>
<td>4</td>
</tr>
</tbody>
</table>
```

**Note:** If the desired assay is not shown, press 7 to display additional assays.
Enter the number for the assay, then press ENTER. All QC test results for the specified assay are printed.

d. To print results for a specific cuvette lot number, press 4 to display the Print by CVlot screen:

```
<table>
<thead>
<tr>
<th>Print by CVlot</th>
</tr>
</thead>
<tbody>
<tr>
<td>ABC...</td>
</tr>
</tbody>
</table>
```

Scan or type the cuvette lot number, then press ENTER. All QC test results for the specified cuvette lot number are printed.

e. To print results for a specific control lot number, press 5 to display the Print by QClot screen:

```
<table>
<thead>
<tr>
<th>Print by QClot</th>
</tr>
</thead>
<tbody>
<tr>
<td>ABC...</td>
</tr>
</tbody>
</table>
```

Scan or type the control lot number, then press ENTER. All QC test results for the specified control lot number are printed.

f. Press CANCEL to return to the menu for printing QC results (Step 1).

**To Check Printer Function:**

1. Press 6 to output an ASCII character set to an attached printer to verify that the printer port functions properly.

```
Hemochron Elite Print Test
!"#$%&'()*+,-./0123456789:;<=>?ABCDEFGHIJKLMNOPQRSTUVWXYZ[\]^_`abcdefgh
ijklmnopqrstuvwxyz[^_`abcdefgh
```

**To Abort Printing of Results:**

1. Pressing CANCEL for three seconds will abort data transmission to the printer. Any data transmitted before CANCEL is released will complete printing.

**Accessing Print Menu-2**

Print Menu-2 allows printing of system information, entering a header, and checking of the printer and barcode scanner.
1. Press PRINT/SCAN to display the first page of the Print menu:

![Print Menu 1]

2. Press 7 to display the second page of the Print menu:

![Print Menu 2]

**To Enter a Custom Header:**
This feature allows creation of a custom header to appear on the top of each printout. The default header is - - -ITC - - -. A custom header can contain up to 20 characters.

1. Display the second page of the Print menu.
2. Press 1 to enter a custom header:

   ![Enter Header]

   ABC

3. Create the custom header using the keypad.
   Press the NOTE key to change the data entry mode from alphabetical (ABC; shown above) to numerical (123) or vice versa.

   **Note:** If alphabetic characters are being entered, advance from the displayed letter to the next letter on that key by pressing the key again before the cursor advances. Release the key (for at least one second) when the selected character is displayed. The character is saved and the cursor advances to the right.

   Use the CANCEL key as a left-arrow key and the ENTER key as a right-arrow key.
4. When the desired header is displayed, press and hold the ENTER key (for at least one second) to save the header. Press and hold the CANCEL key to return to the ENTER HEADER prompt without saving the new header.

**To Print System Information:**

1. Display the second page of the Print menu.
2. Press 2 to print the system software version, system serial number, and program parameters.

**Other Database Operations**
Results can be reviewed for both patient and QC tests. The total number of test results that are currently stored can be displayed, the total number of additional test results that can be stored can be displayed, and the results in either the patient database or the quality control database can be erased to make room for additional test results. Finally, POCT-1A communication can be initiated.

**CAUTION:** Results from the oldest test are overwritten if a test is run when the database is full. Therefore, it is important to periodically check, print, archive, and erase database contents.

**To Perform Other Database Operations:**
1. Press DATABASE to display the Database menu:

   ![Database]

   1-Totals
   2-Pat Erase
   3-QC Erase
   4-Display Pat
   5-Display QC
   6-ITC >> COM
Then perform one of the following procedures:

**To Display Total Tests and Remaining Tests in the Database:**
1. Press 1. The total number of stored patient records and QC records, plus the number of records that can still be added to the patient database and the QC database are displayed:

<table>
<thead>
<tr>
<th>Pat DB Totals</th>
<th>QC DB Totals</th>
</tr>
</thead>
<tbody>
<tr>
<td>122 Tests/488 Left</td>
<td>24 Tests/576 Left</td>
</tr>
</tbody>
</table>

**To Delete All Results from a Database:**

*Note*: Deleting of results from the database can be disallowed in Configuration Manager. Refer to page 53 for details.
1. Press 2 to erase the patient database.
   - Or, press 3 to erase the QC database.
2. The instrument displays a prompt to confirm deletion of the database:

   Sure? 1-YES 2-NO

3. Press 1 to confirm deletion. COMPLETED is displayed for two seconds.

   *Note*: Press 2 or CANCEL to cancel erasing of the database.

**To Review Results of the Patient Database:**
1. Press 4. The first page of the results for the most current test record is displayed.

2. Press 9 to display additional pages of the record:

3. To scroll through additional test records, press ENTER (for the next record) or 0 (for the previous record).

**To Review Results of the QC Database:**
1. Press 5. The first page of the results for the most current test record is displayed.

2. Press 9 to display additional pages of the record:

3. To scroll through additional test records, press ENTER (for the next record) or 0 (for the previous record).

**To Initiate Communication When Using the POCT-1A Communication Protocol:**
1. Press 6. A confirmation is displayed when communication is established:
To Exit Database Operations:

1. Press CANCEL to exit.

Downloading QC/Patient Data to Your PC

For information on printing QC/Patient data through your PC using Accriva Diagnostics, Inc. data management software, contact Technical Support.

Acronyms for Error Messages

The following table lists the acronyms for Error Messages when viewing stored Patient or QC records:

<table>
<thead>
<tr>
<th>Acronym</th>
<th>Description</th>
<th>Display Message</th>
</tr>
</thead>
<tbody>
<tr>
<td>ORH</td>
<td>Out of Range High</td>
<td>Out of Range-Hi</td>
</tr>
<tr>
<td>ORL</td>
<td>Out of Range Low</td>
<td>Out of Range-Lo</td>
</tr>
<tr>
<td>SNS</td>
<td>Sample Not Seen</td>
<td>Sample Not Seen</td>
</tr>
<tr>
<td>STS</td>
<td>Sample Too Small</td>
<td>Sample Too Small</td>
</tr>
<tr>
<td>STL</td>
<td>Sample Too Large</td>
<td>Sample Too Large</td>
</tr>
<tr>
<td>TTO</td>
<td>Temp Timeout</td>
<td>Heater Too Cool</td>
</tr>
<tr>
<td>TTH</td>
<td>Temp Too High</td>
<td>Heater Too Hot</td>
</tr>
<tr>
<td>SPF</td>
<td>Sample Position Fault</td>
<td>Sample Pos Fault</td>
</tr>
<tr>
<td>UAT</td>
<td>User Aborted Test</td>
<td>User Abort</td>
</tr>
<tr>
<td>BAT</td>
<td>Battery Fault</td>
<td>Battery Fault</td>
</tr>
<tr>
<td>PMS</td>
<td>Premature Sample</td>
<td>Premature Sample</td>
</tr>
<tr>
<td>DEF</td>
<td>Detector Fault</td>
<td>Detector Fault</td>
</tr>
<tr>
<td>BTH</td>
<td>Battery Too High</td>
<td>Battery Too High</td>
</tr>
<tr>
<td>XTH</td>
<td>External Power Too High</td>
<td>External Too High</td>
</tr>
<tr>
<td>RTC</td>
<td>Real Time Clock Fault</td>
<td>RTC Fault</td>
</tr>
</tbody>
</table>
COMMUNICATION WITH A PC OR NETWORK

The instrument can communicate directly with a personal computer (using the COM port) or a network (using the Ethernet port). One of two communication protocols can be selected:

- Accriva proprietary communication protocol that allows records to be transferred from the instrument and allows the instrument to be configured using software such as Hemochron Configuration Manager.
- A communication protocol that meets CLSI POCT-1A standards. This protocol allows the instrument to communicate with any software or device that is compliant with POCT-1A.

**Note:** The POCT-1A communication protocol is only allowed when Port is set to NET.

Communicating When the POCT-1A Communication Protocol is Used

When the CLSI POCT-1A compliant communication protocol is used, communication must be initiated from the instrument.

1. If necessary, check the instrument Communications menu and verify that the POCT-1A communication protocol and the NET communication port are specified:

   ![Communications Menu](image)

   **Note:** The Communications menu is accessed from the Supervisor menu (page 21).

2. From either the Database menu or the Communications menu, select 6–POCT >> NET to initiate communication:

3. A confirmation is displayed when communication is established:

4. At the software application (for example, Hemochron Configuration Manager on the personal computer), start data transfer. Progress of data transfer is displayed as data transfer proceeds:

   - **Example:** Reading the configuration from an instrument using Hemochron Configuration Manager software.
   - **Example:** Writing a new configuration to an instrument using Hemochron Configuration Manager software.
   - **Example:** Exporting test records from an instrument using Hemochron ReportMaker software.

5. The message Disconnecting Network is displayed upon completion of data transfer, and the instrument main menu is then displayed.
Communicating When the Proprietary Communication Protocol is Used

When the proprietary communication protocol is used, communication must be initiated from the personal computer or network.

1. If necessary, check the instrument Communications menu and verify that the proprietary communication protocol and the desired communication port (NET or COM) are specified:

<table>
<thead>
<tr>
<th>Communications</th>
<th>Communications</th>
</tr>
</thead>
<tbody>
<tr>
<td>1&gt;=POCT OFF</td>
<td>1&gt;=POCT OFF</td>
</tr>
<tr>
<td>2-&gt;Port NET</td>
<td>2-&gt;Port COM</td>
</tr>
<tr>
<td>3-&gt;Push POCT OFF</td>
<td>3-&gt;Push POCT OFF</td>
</tr>
<tr>
<td>4-&gt;PORT Setup</td>
<td>4-&gt;PORT Setup</td>
</tr>
<tr>
<td>5-&gt;XPORT COM1-SET</td>
<td>5-&gt;XPORT COM1-SET</td>
</tr>
<tr>
<td>6-&gt;ITC &gt;&gt; NET</td>
<td>6-&gt;ITC &gt;&gt; COM</td>
</tr>
</tbody>
</table>

*Note: The Communications menu is accessed from the Supervisor menu (page 21).*

2. At the software application (for example, Hemochron Configuration Manager on the personal computer), start data transfer. Status messages may be displayed on the instrument during initiation of communication:

- XPORT Power-up
- Connecting
- ITC >> NET
- CANCEL to Exit

and confirmation is displayed when communication is established:

3. Initiate data transfer at the software application. Confirmation of data transfer is displayed as data transfer proceeds:

- Transferring
- Data

4. The message *Shutting Down* is displayed upon completion of data transfer, and the instrument then shuts down.

Notification of Idle Network

A message on the instrument alerts the operator if the network is idle for more than five seconds during a communication session:

- Network
- Idle
- CANCEL to Exit

Cancelling a Communication Session

A communication session can be cancelled by pressing the CANCEL key on the instrument. This can be done whenever the CANCEL prompt is displayed on the instrument:

- ITC >> NET
- CANCEL to Exit
CONFIGURATION MANAGER

Hemochron Configuration Manager (HCM) software V3.0 or higher allows the supervisor to configure Hemochron Signature Elite instruments so they meet the needs of the laboratory. The configuration is performed on a personal computer with installed Hemochron Configuration Manager software. The computer is connected to the instrument by means of a COM port or a network connection. The personal computer must meet the following minimum requirements:

- Pentium II 400 MHz or higher microprocessor
- Hard disk with at least 50 MB of free space
- At least 128 megabytes (MB) of random-access memory (RAM)
- Microsoft Windows NT, 2000, or XP (if you are running Windows 2000 or XP, you must have System Administrator security privileges to install the software)
- VGA 800x600 or higher resolution supported by Microsoft Windows.
- At least one available RS-232 serial port (if an instrument is connected directly to the personal computer by means of a COM port)
- A network connection (if an instrument is connected to the personal computer by means of a network)
- CD-ROM drive

Installing Configuration Manager Software

The Setup program creates Hemochron Configuration Manager working directories. It copies programs and configuration files to the working directories and adds the Hemochron Configuration Manager (CM) program icon to the desktop.

1. Start Microsoft Windows.
2. Insert the Hemochron Configuration Manager installation CD-ROM in the CD drive.
3. Follow the instructions on the screen.
   
   **Note:** You will be prompted to specify a folder for the installation.
4. A message will be displayed when the software is successfully loaded. After the message is displayed, remove the CD from the computer.
5. Shut down and restart the computer if prompted to do so.

Connecting an Instrument

The Hemochron Signature Elite instrument that is being configured must be connected to Hemochron Configuration Manager. This can be done using either the COM port or a network connection.

1. Connect the Hemochron Signature Elite instrument to the personal computer on which Hemochron Configuration Manager is installed, using either the computer interface cable included with the instrument or a network interface cable.
   
   **Note:** If the computer interface cable is used, connect one end of the cable to the computer port on the side of the instrument and the other end to an unused COM port on the personal computer. If a network interface cable is used, connect one end of the cable to the Ethernet port on the side of the Hemochron Signature Elite instrument and the other end to a network connection port. Refer to page 24 for connection details.
   
   **Note:** A computer interface cable is included with the instrument and CD, or a separate PCKIT (Personal Computer Interface Cable) can be obtained from Customer Service. Alternatively, the cable can be prepared as outlined on page 25.
   
   **Note:** It is recommended that the instrument is on external power while it is connected to the personal computer.
2. Check the instrument Communications menu and verify that the desired communication protocol and communication port (NET or COM) are specified (page 21).
   
   **Note:** When the POCT-1A communication protocol is used, Hemochron Configuration Manager cannot communicate with the instrument over the COM port.
3. Follow instructions on page 44 for establishing communication.
Starting Configuration Manager

1. Select the Accriva Diagnostics, Inc. CM program icon. The Startup window is displayed:

   ![Configuration Manager Startup Window]

   - Click OK to logon using the default user name and password:
   - Click OK to start the Configuration Manager.

2. When Hemochron Configuration Manager software is run for the first time, instructions are displayed to logon using the default user name and password:

   ![Logon Instructions]

   a. Click OK and then click Start Program. The Logon dialog box is displayed with the default user name (supervisor) and password (supervisor):

   ![Logon Dialog Box]

   b. Click OK. The System Configuration dialog box is displayed:
c. Enter the institution name, address, and/or contact information.
   Click **Auto-Detect** to automatically detect the PC COM port that is being used when connected to *Signature Elite*. Alternately, click the drop-down arrow in the COM Port box and select the number of the COM port used.
   **Note:** To modify these settings at a later date, select **System Configuration** from the Utilities menu.

d. Click **OK**. The information is saved, and the User Master File Maintenance dialog box is displayed:

e. To change the default “Supervisor” user password, enter a new password. Ensure that the **Yes** button is selected for Supervisor access, then click **Save**.
   **Note:** The “Supervisor” user name cannot be deleted. However, the default user password should be changed as shown above. All commands are accessible to anyone until then.

3. Thereafter, click **Start Program** to display the Logon dialog box:

4. Enter your user name and password, then click **OK**.

**Menus**

Menus for Hemochron *Configuration Manager* functions are displayed at the top of the main window:
The use of each menu is described below:

<table>
<thead>
<tr>
<th>Menu</th>
<th>Purpose</th>
</tr>
</thead>
<tbody>
<tr>
<td>Configuration</td>
<td>Display the Read Configuration Data dialog box. This dialog box allows the operator to set the various configuration options (see page 52).</td>
</tr>
<tr>
<td>Reports</td>
<td>Display and print listings of configuration parameters, instruments, operators (see page 65), and audit trail.</td>
</tr>
<tr>
<td>Master File Maintenance</td>
<td>Display, modify, add, and delete instrument information.</td>
</tr>
<tr>
<td>Utilities</td>
<td>Rebuild and compress the database. Purge selected records. Set up system time check. Display, modify, add, and delete operator information. Convert Hemochron Configuration Manager software V1.0 or V2.0 data to V3.0.</td>
</tr>
<tr>
<td>Exit</td>
<td>Log off and/or exit the system.</td>
</tr>
<tr>
<td>Help</td>
<td>Display the version number of Hemochron Configuration Manager software that is installed. Access Help.</td>
</tr>
</tbody>
</table>

**Reading or Loading Configuration Data**

Select **Configuration** in the **Configuration Manager** main menu to display the Read Configuration Data dialog box. This dialog box allows configuration data to be read from a connected instrument or a different configuration file to be loaded:
Note: Hemochron Configuration Manager uses a Microsoft Windows user interface. Software features are intuitive and easy to learn. Refer to Windows documentation for instructions on navigation, access to commands and Help, execution of commands, and shortcuts that can be taken.

Note: The Instrument Type drop-down list is used when loading a configuration from the database. If a configuration is read from a connected instrument, the instrument type is automatically detected and displayed.

Note: Load the default Hemochron Signature Elite settings by selecting Factory Default from the Configuration/Revision drop-down list. These settings can be modified and saved under a different configuration file name. The Factory Default configuration file cannot be overwritten or purged from the database.

To Read Configuration Data from a Connected Instrument:

1. Click Read From Instrument. The Read Configuration from Instrument dialog box is displayed:

2. Establish communication to the instrument as described on pages 44 and 45.

   Note: If the instrument is connected over a network, select <Listen on Network> from the drop-down list in the Connect via box. Otherwise, if prompted, designate the COM port by selecting it from the drop-down list.

   Note: When the POCT-1A communication protocol is used, Hemochron Configuration Manager cannot communicate with the instrument over the COM port.
3. Click **Start**. Configuration data are read from the connected instrument:

**Note:** The instrument serial number and the name and revision level of the instrument configuration file are displayed in the Read Configuration dialog box. Progress of reading of configuration data is displayed in the status lines.

**Note:** If the proprietary communication protocol is used, **Transferring Data** is displayed on the connected Hemochron Signature Elite instrument during data transfer. If the connection is not made, an error message is displayed on the personal computer. Select **Retry** on the error message box to make the connection.

**Note:** If the POCT-1A communication protocol is used, specific messages indicate the progress of data transfer. If the connection is not made, the instrument will beep, display **Network Timed Out!**, and prompt that the instrument is ready to accept a cuvette.

4. When reading of configuration data is successfully completed, a confirmation dialog box is displayed. Click **OK** to display the Configuration Module dialog box (see below).

**To Load a Configuration File from the Database:**
1. Select the instrument type from the **Instrument Type** drop-down list.
2. Select the file from the **Select Existing Configuration** drop-down list.
3. Click **Yes** to the prompt for loading the selected file.
4. Click **OK** when reading of the configuration data is complete. The Configuration Module dialog box is displayed:
### Setting Configuration Options

Reading a configuration file from an instrument or loading a configuration file from the database will open the Configuration Module dialog box (page 51). Buttons on the dialog box allow the supervisor to display and/or modify the instrument configuration:

<table>
<thead>
<tr>
<th>Button</th>
<th>Purpose</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Instrument Options</strong></td>
<td></td>
</tr>
</tbody>
</table>
| **a. General Options** | 1. Specify when the display panel is illuminated.  
2. Allow operators to change the date or time.  
3. Allow operators to erase a database.  
4. Suppress display of QC results.  
5. Enter instrument supervisor PIN. |
| **b. Operator ID Options** | 1. Require entry of an Operator ID (OID) before a test can be run.  
2. Require that the supervisor-specified OID or Personal Identification Number (PIN) must be entered by an operator in order to operate the instrument.  
3. Specify that the OID for the previous test is automatically entered for a specified length of time or until a new OID is entered.  
4. Specify that the barcode scanner must be used for entering the Operator ID. |
| **c. Patient ID Options** | 1. Specify the number of digits for a Patient ID (PID).  
2. Require entry of a (PID) before a test can be run.  
3. Specify that the PID for the previous test is automatically entered for a specified length of time or until a new PID is entered.  
4. Specify that the barcode scanner must be used for entering the Patient ID. |
| **d. Date Format Options** | 1. Specify a date format of MM/DD/YY or YY-MM-DD. |
| **e. Time Format Options** | 1. Specify a **24-hour** or **12-hour** (am and pm) time format. |
| **f. Cuvette Lot Options/QC Lot Options** | 1. Require entry of a cuvette lot number and/or QC lot number.  
2. Specify that the barcode scanner must be used for entering the cuvette lot number and/or QC lot number.  
3. Require that cuvette and/or QC lot numbers that are used can only be selected from a list specified by the supervisor. |
| **Lock-Out Control** | |
| **a. QC Lockout** | 1. Specify whether electronic quality control (EQC) testing must be run after a specified time interval has elapsed.  
2. Specify either 300 seconds or 500 seconds for the EQC level 2 test.  
3. Specify automatic running of EQC tests at scheduled intervals.  
4. For each test type, specify whether normal and/or abnormal liquid quality control (LQC) testing must be run after a specified time interval has elapsed.  
5. Prohibit specific test(s) from being performed on the instrument.  
6. If EQC and/or LQC testing must be run after a specified interval, specify the number of additional tests that can be run per assay (the 911 exceptions) after the time interval has elapsed. |
<p>| <strong>b. QC Exceptions</strong> | 1. Specify the number of EQC or LQC test failures that are allowed before the instrument is locked and must be reset by the supervisor. |</p>
<table>
<thead>
<tr>
<th>Button</th>
<th>Purpose</th>
</tr>
</thead>
<tbody>
<tr>
<td>Operator Table</td>
<td>1. Specify an OID, PIN, and/or certification expiration date for an operator.</td>
</tr>
<tr>
<td>User Notes</td>
<td>1. Specify user-selectable notes that can be entered at the time of a test.</td>
</tr>
<tr>
<td>Cuvette Lots</td>
<td>1. Specify up to five cuvette lot numbers and expiration dates that are to be used for each assay.</td>
</tr>
<tr>
<td>QC Lots</td>
<td>1. Specify up five lot numbers, expiration dates, and expected ranges for level 1 (normal) and level 2 (abnormal) controls for each assay.</td>
</tr>
</tbody>
</table>

**Specifying Instrument Options**

Instrument options specify how the instrument communicates with the operator.

1. Click **Instrument Options** on the Configuration Module dialog box (page 51) to display the Instrument Options dialog box. Information concerning the configuration file is displayed:

![Instrument Options (SignatureElite)](image)

### Instrument General Options

1. **Allow Date & Time Change**: Enter a check mark to allow the operator to change the date and time on the instrument.
2. **Allow Patient & QC Database Erase**: Enter a check mark to allow an operator to erase the patient database or the QC database.
3. **LQC Hide**: Enter a check mark to specify that Pass or Fail is displayed in place of numerical QC Results.
4. **Supervisor PIN**: Enter an optional PIN number (from 1 to 9 digits) to control access to the Supervisor menu in the Hemochron Signature Elite instrument.
5. **Flashlight**: Enter the number of minutes that the display will be illuminated after a cuvette is inserted and after results are displayed. Valid range for screen illumination is from 0 to 100 minutes.

*Note: If 0 is entered, the display is illuminated at all times.*
Operator ID Options

1. **Not Required**: Select if entry of an Operator ID (OID) or PIN is not required when running a test.
2. **OID Required**: Select to require entry of an OID (any OID) when running a test.
3. **Valid OID Required**: Select to require entry of an OID that is included in the Operator Table and is within the expiration period (if an expiration date is enabled for that user).
4. **Valid PIN Required**: Select to require entry of an authorized Operator PIN. All PIN numbers must have a corresponding Operator ID in the Operator Table.
   
   *Note: Only one of the alternatives listed in steps 1 through 4 can be selected. Authorized OIDs and/or PINs are entered in the Operator Table. Both OID and PIN should be unique for each authorized operator or group in the Operator Table (see page 59).*

5. **OID Scan Only**: Enter a check mark to specify that the OID must be scanned.
   
   *Note: If OID Scan Only is enabled, the operator is prompted to scan the OID when running a test. The operator can still enter the OID manually by pressing the ID key (as indicated by the ID=Manual Entry prompt), but the OID must be entered twice.*

6. **Reuse Operator ID**: Enter a value (from 1 to 240 hours) to allow the instrument to use the OID from the previous test as the default entry for the current test. The OID will be entered by default for the specified length of time and will be flagged as a manual entry in the database.
   
   *Note: If 0 is entered, the OID will not be reused. The default OID can be changed if needed.*

Patient ID Options

1. **Require Patient ID**: Enter a check mark to require entry of a PID when running a patient test.
2. **Patient ID Length**: Select the minimum (from 1 to 20) and maximum (from 1 to 20) number of digits that must be entered for a patient ID.
3. **PID Scan Only**: Enter a check mark to specify that the PID must be scanned.
   
   *Note: If PID Scan Only is enabled, the operator is prompted to scan the PID when running a test. The operator can still enter the PID manually by pressing the ID key (as indicated by the ID=Manual Entry prompt), but the PID must be entered twice.*

4. **Reuse Patient ID**: Enter a value (from 1 to 240 hours) to allow the instrument to use the PID from the previous test as the default entry for the current test. The PID will be entered by default for the specified length of time and will be flagged as a manual entry in the database.
   
   *Note: If 0 is entered, the PID will not be reused. The default PID can be changed by entering a new PID.*

Date Format

1. **Date Format**: Select MM/DD/YY to display June 1, 2006 as 06/01/06. Select YY-MM-DD to display June 1, 2006 as 06-06-01.

Time Format

1. **Time Format**: Select 24 Hour to display 7:15 pm as 19:15. Select 12 Hour to display 7:15 pm as 7:15 pm.

Cuvette Lot Options

1. **Required**: Enter a check mark to require entry of a cuvette lot number and expiration date when running a test.
2. **Scan only**: Enter a check mark to specify the cuvette information can only be entered using the barcode scanner.
   
   *Note: If Scan Only is enabled, the operator is prompted to scan the cuvette information when running a test. The operator can still enter the information manually by pressing the 7 key (as indicated by the 7=Manual Entry prompt), but the information must be entered twice. Refer to page 28 for additional information.*
3. **Pre-stored only**: Enter a check mark to specify the cuvette information can only be selected from a list of lot numbers specified by the supervisor.
   
   **Note**: Up to five cuvette lot numbers and expiration dates can be specified for each assay (see page 61).

   **Note**: If more than five cuvette lot numbers per assay are scanned or entered, the lot number(s) with the closest expiration date will be deleted.

**QC Lot Options**

1. **Required**: Enter a check mark to require entry of a QC lot number when running a quality control test.
2. **Scan only**: Enter a check mark to specify the QC lot number can only be entered using the barcode scanner.
   
   **Note**: If **Scan only** is enabled, the operator is prompted to scan the QC information when running a test. The operator can still enter the QC information manually by pressing the 7 key (as indicated by the **7=Manual Entry** prompt), but the QC information must be entered twice. Refer to page 32 for additional information.

3. **Pre-stored only**: Enter a check mark to specify the QC lot number can only be selected from a stored list.
   
   **Note**: Up to five QC lot numbers, expiration dates, and ranges can be specified for each assay for both level 1 and level 2 controls (see page 62).

   **Note**: If more than five QC lot numbers per assay are scanned or entered for level 1 and/or level 2 controls, the lot number(s) with the closest expiration date will be deleted.

**Saving the Options**

1. Click **Save & Exit** to save the specified options and cancel display of the dialog box.
   
   **Note**: **Cancel** clears all fields without saving any parameters.

**Specifying QC Lockout Parameters**

QC lockout parameters specify the maximum length of time that can elapse before liquid quality control (LQC) or electronic quality control (EQC) must be successfully run (**QC Interval**). When the lockout interval has elapsed and the test cuvette is inserted into the instrument, the user will be prompted to perform the appropriate QC.

If 911 Limits are enabled, the user will be offered the choice to run a patient test. If this option is selected, the instrument displays the number (from 0 to 240) of remaining tests per assay that can be run (the 911 override) before QC must be performed (see **To Specify QC Exceptions** on page 58). Separate requirements can be specified for the frequency at which EQC and LQC for various tests must be run at normal and abnormal levels.

**To Specify Lockout Parameters for All Tests Except EQC**:

1. Click **Lock-Out Control** on the Configuration Module dialog box (page 51) to display the QC Lockout dialog box. Select the QC Lock Out Requirements tab (if needed).
2. Select the test for specifying lockout parameters. Current settings for the test are displayed:
Note: The number of patient tests that have already been run against the 911 override limit is displayed in the # of 911s Used field when a configuration is read from an instrument. This value is automatically reset to zero when QC is successfully run.

3. Enter a check mark in the **QC Normal Required** box to require QC Normal testing to be run at a specified interval.

   Note: The number of QC Normal tests that have been run and failed is displayed in the # of QC Tests Failed field.

4. (If QC Lot Options are not specified) Designate the acceptable range for Normal LQC of the specified test by entering a value in the **Low** and **High** fields.

   Note: If **Required, Scan only**, and/or **Pre-stored only** is specified for QC Lot Options, the **Low** and **High** fields are not displayed.

5. Enter a check mark in the **QC Abnormal Required** box to require QC Abnormal testing to be run at a specified interval.

   Note: The number of QC Abnormal tests that have been run and failed is displayed in the # of QC Tests Failed field.

6. (If QC Lot Options are not specified) Designate the acceptable range for Abnormal LQC of the specified test by entering a value in the **Low** and **High** fields.

   Note: If **Required, Scan only**, and/or **Pre-stored only** is specified for QC Lot Options, the **Low** and **High** fields are not displayed.

7. (If the QC testing displayed on the tab is required) Enter the maximum length of time (the QC Interval, from 0 to 1080 hours) that can elapse before the QC testing must be successfully run.

   Note: The interval time resets upon successful completion of QC.

8. (If the QC testing displayed on the tab is required) Enter the date and time when the required QC testing displayed on the tab must be run for the first time. If a start date and time are not entered, the QC testing will be required immediately after Configuration is downloaded to an instrument.

   Note: Subsequent dates and times will be determined automatically from the QC interval.

9. Repeat steps 2 through 6 as needed for additional QC testing.

10. Click **Save** to save the QC lockout parameters for each test type individually.

    Note: **Click Cancel** to clear the fields without saving the modified parameters. **Save & Exit** saves the EQC/LQC lockout parameters and closes the dialog box.

11. To disable a particular assay from being run, enter a check mark in the **QC Normal** (or **Abnormal** Required) box, enter a range, and then enter 0 for the QC Interval. The
message
DISALLOWED ASSAY … REMOVE CUVETTE displays when a cuvette for that test is inserted.

To Specify Lockout Parameters for EQC:

1. Click Lock-Out Control on the Configuration Module dialog box (page 51) to display the QC Lockout dialog box. Select the QC Lock Out Requirements tab (if needed), and then select EQC:

   ![EQC Lockout Dialog Box](image)

2. Enter a check mark in the Required box to specify that EQC testing is to be run at a specified interval.
   
   **Note:** The number of EQC tests that have been run and failed is displayed in the # of QC Tests Failed fields. The expected range for each level is displayed in the Low and High fields.

3. Select either 300 seconds or 500 seconds for the level 2 EQC test.
4. (If EQC testing is required) Enter the maximum length of time (the QC Interval, from 1 to 1080 hours) that can elapse before EQC testing must be successfully run.
   
   **Note:** The interval time resets upon successful completion of EQC.

5. (If EQC testing is required) Enter the date and time when EQC testing must be run for the first time. If a start date and time are not entered, EQC testing will be required immediately after the configuration is downloaded to an instrument.
   
   **Note:** Subsequent dates and times will be determined automatically from the QC interval.

6. **Auto EQC:** Enter a check mark to specify that EQC should be automatically run at the specified interval.
   
   **Note:** If Auto EQC is selected and the Hemochron Signature Elite is connected to a power adapter, the instrument will remain on and perform the required EQC test at the appropriate interval. The results will be automatically stored in the database.
   
   **Important:** If Auto EQC is selected, ensure that cuvettes are routinely removed from the instrument after completing a test. If EQC is run while a cuvette is inserted, the message **Detector Blocked** is displayed and the EQC test fails.

7. Click Save to save the EQC lockout parameters.
   
   **Note:** Click Cancel to clear the fields without saving the modified parameters. Save & Exit saves the EQC lockout parameters and closes the dialog box.

To Specify QC Exceptions:
1. Click **Lock-Out Control** on the Configuration Module dialog box (page 51) to display the QC Lockout dialog box.
2. Select the QC Exceptions tab:

![QC Lockout dialog box](image)

**Note:** Entries made on the QC Exceptions tab apply to all assays.

### 911 Limit
1. Enter the maximum number of patient tests (the 911 Limit, from 0 to 240 tests) that can be run after the specified EQC/LQC lockout interval has been reached an assay. When the 911 limit is reached for an assay, that assay cannot be run until a successful QC test is run for that assay or until the instrument is unlocked (see page 58) using Configuration Manager.

   **Note:** The number of patient tests that have already been run against the current 911 limit is displayed when read from an instrument.

### Failure Limit
1. EQC Failure Limit: Enter the maximum number of times that a failed EQC test can be rerun before the system will lock (from 1 to 9 tests).
2. LQC Failure Limit: Enter the maximum number of times that a failed LQC test can be rerun before the system will lock (from 1 to 9 tests).

   **Note:** Setting the failure limit to 0 (zero) disables lockout by failure count.

### Saving the Options
1. Click **Save** to save the QC Exceptions parameters.

   **Note:** **Cancel** clears all fields without saving any parameters. **Save & Exit** saves the EQC/LQC lockout parameters and closes the dialog box.

### To Unlock an Instrument that has Reached QC Failure Limit or the 911 Limit

**Note:** The Hemochron Signature Elite must be connected to the Configuration Manager and the instrument configuration information read by clicking on **Read Configuration from Instrument**.

1. After the configuration data from the instrument has been read, select **Clone Instrument**. The Clear Instrument Counts dialog box is displayed (see page 63).
2. Place check marks in both check boxes to reset the failure counts and 911 use counts on cloning.
3. Click **OK**. The Write Configuration dialog box is displayed (page 63).
4. Click **Start**. The reset configuration is written to the instrument.
Note: A message box is displayed when writing of the configuration data is complete. Click OK to acknowledge the message. Then click Exit to cancel display of the Configuration Module dialog box.

Specifying Operator IDs and/or Personal Identification Numbers

The Operator Table allows the supervisor to enter a name for each operator and then specify an Operator ID (OID), Personal Identification Number (PIN), and/or certification expiration date for each operator (or a group of operators). The supervisor can then specify in the Instrument Options dialog box (see page 53) whether entry of an OID or PIN that is included on the Operator Table is required before the instrument can be operated.

1. Click Operator Table on the Configuration Module dialog box (page 51) to display the Operator Table dialog box:

2. (Optional) Enter the name of each operator.
   Note: The operator’s name is for reference only. It is not stored in the instrument.

3. Enter or scan an Operator ID (up to 16 alphanumeric characters) for each operator.
   Note: OIDs may be scanned using the barcode scanner through the Supervisor menu and uploaded to Configuration Manager.
   Note: If OIDs will be entered using the instrument keypad, OIDs should contain only those letters, numerals, and/or characters that can be entered using the 0 through 9 keys on the keypad.
   Note: An OID that is not included in the Operator Table will not be allowed to run a test if Valid OID Required is selected in the Instrument Options dialog box (see page 53). If OID Required is selected, any OID can be entered when running a test.

4. (Optional) Enter a numeric PIN (up to 9 digits) for each operator. The PIN can be any number other than zero (0).
   Note: A unique PIN can be assigned for each authorized operator, or a single OID and PIN can be used to allow operation by any operator that enters the correct code.
5. (Optional) Specify an expiration date for certification of an operator by entering a check mark in the Expires box for that operator and then selecting the expiration date from the pop-up calendar.

   **Note:** An operator with expired certification will not be allowed to run a test if Valid OID Required is selected in the Instrument Options dialog box (see page 53). If OID Required is selected, any OID can be entered when running a test.

6. To view OIDs without a PIN or expiration date, select the drop-down arrow in the Sort By box and select PIN or Expiration Date to sort the OIDs in order of their PIN or expiration date. This places OIDs without a PIN or expiration date at the beginning of the table.

   **Important:** Sorting OIDs by expiration date places those OIDs with the expiration field unchecked and grayed at the beginning of the list. However, the dates that are displayed in the grayed fields are not refreshed and are incorrect. Expiration dates that are displayed for OIDs with enabled (checked) expiration dates are correct.

7. The OID/PIN Requirement box displays the OID/PIN requirements that are currently set under Instrument Options (see page 53). These options are all mutually exclusive.

8. Click Save & Exit to save changes to the Operator Table and exit the dialog box.

   **Note:** Cancel removes the dialog box without saving changes. Clear Current Page clears all information from the current page of the Operator Table. Clear All clears all information from the entire Operator Table.

**Specifying User Notes**

Up to nine customized notes can be entered into the User Notes menu of an instrument. Up to two of these notes can be selected and appended to a test record by the operator when the test is run.

1. Click User Notes on the Configuration Module dialog box (page 51) to display the User Notes dialog box. Each field label includes the instrument numeral key that will enter that note when the User Notes menu is displayed on the instrument (page 33):

2. Enter a note (up to 16 characters) for each key.

   **Note:** No note is displayed on the Hemochron Signature Elite instrument if a note is not entered for that note number.

3. Click Save & Exit to save the specified notes and exit the dialog box.

   **Note:** Reset clears all fields. Cancel cancels display of the dialog box without saving the modified notes.
Entering Cuvette Lot Numbers and Expiration Dates
Up to five cuvette lot numbers and expiration dates can be entered for each assay type. Selection or scanning of a cuvette lot number is only allowed when Required, Scan only, and/or Pre-stored only is specified for Cuvette Lot Options (see page 54).

*Note:* Cuvette lot numbers may be scanned using the barcode scanner through the Supervisor menu and uploaded to Configuration Manager (page 18).

*Note:* Cuvette lot numbers that are manually entered into the Hemochron Signature Elite instrument when running a test are only saved to that particular test record and are not uploaded to Configuration Manager.

1. Click **Cuvette Lots** on the Configuration Module dialog box (page 51) to display the Cuvette Lot Number Entry dialog box:

2. Select an assay from the **Select Test Type** drop-down list to display the form for entering cuvette lot numbers for that assay and their expiration dates:

3. Enter the lot number and expiration date for up to five cuvette lots.
4. Click **Save** to save the cuvette lot information for the specified assay.

*Note:* **Cancel** clears and closes the Cuvette Lot Number Entry dialog box without saving entries. **Exit** closes the Cuvette Lot Number Entry dialog box.
Entering QC Lots

Up to five quality control lot numbers, expiration dates, and expected ranges can be entered per assay type for both level 1 (normal) and level 2 (abnormal) liquid controls. Selection or scanning of a QC lot number is only allowed when Required, Scan only, and/or Pre-stored only is specified for QC Lot Options (see page 55).

Note: If more than five QC lot numbers per assay are scanned or entered for level 1 and/or level 2 controls, the lot number(s) with the closest expiration date will be deleted.

Note: QC lot number information may be scanned using the barcode scanner through the Supervisor menu and uploaded to Configuration Manager (page 18). If needed, the expected ranges can then be manually changed by the supervisor in Configuration Manager.

Note: QC lot number information that is manually entered into the Hemochron Signature Elite instrument when running a QC test is only saved to that particular QC test record and is not uploaded to Configuration Manager.

1. Click QC Lots on the Configuration Module dialog box (page 51) to display the QC Lot Number Entry dialog box:

2. Select an assay from the Select Test Type drop-down list to display the form for entering QC lot numbers, expiration dates, and expected ranges for that assay:
3. Enter the QC lot number, expiration date, and low and high limits for level 1 (normal) and/or level 2 (abnormal) controls. The limits that are entered may be from the expected range contained in the labeling for the controls or from an expected range specified by the supervisor.

4. Click **Save** to save the QC lot number information for the specified assay.

   **Note:** **Cancel** clears and closes the QC Lot Number Entry dialog box without saving entries. **Exit** closes the QC Lot Number Entry dialog box.

**Saving Modified Configuration Data**

After configuration data has been modified, the data must be saved before it can be downloaded to an instrument or used in the future.

1. Click the **Save Configuration** button on the Configuration Module dialog box (page 51) to display the Save Configuration dialog box:

2. Enter a file name and/or revision for the configuration data to be saved:
   - To save the configuration data under a new file name, enter a name for the new configuration file in the **Configuration** field and enter the revision number for the file in the **Revision** field.
   - To save the configuration data as a new revision of an existing file name, select the existing file name and revision from the **Configuration/Revision** field and enter the new revision number for the file in the **Revision** field.
   - To overwrite an existing configuration file, select the existing file name and revision from the **Configuration/Revision** field.

3. (Optional) Enter information to describe the new configuration in the **Description 1** through **Description 5** fields.

4. Click **OK** to save the configuration, then click **OK** to respond to the confirmation prompt.

   **Note:** If an existing configuration file is being overwritten, a confirmation to overwrite the file is displayed. Click **Yes** to overwrite the file or click **No** to cancel the operation.

**Writing Configuration Data to an Instrument**

After configuration data have been modified and saved, the data from the configuration file (or any other saved configuration file) can be downloaded to a Hemochron Signature Elite instrument.

To **Write Configuration Data to an Instrument**:

1. (If needed) Connect the instrument to the PC (see page 24).
2. (If needed) Start Hemochron **Configuration Manager** and select the configuration file that is to be downloaded (see page 49).
3. Click the **Clone Instrument** button on the Configuration Module dialog box (page 51) to display the Clear Instrument Counts dialog box:

   ![Clear Instrument Counts dialog box](image)

   **Note:** The Clear Instrument Counts dialog box allows the user to specify whether failure counts and 911 use counts will be reset on cloning and to synchronize the instrument time and date with the PC time and date when using the POCT-1A communication protocol.

4. Click **OK** to continue.

5. The Write Configuration dialog box is displayed:

   ![Write Configuration dialog box](image)

6. Establish communication to the instrument as described on pages 44 and 45.

   **Note:** If the instrument is connected over a network, select `<Listen on Network>` from the drop-down list in the Connect via box. Otherwise, if prompted, designate the COM port by selecting it from the drop-down list.

7. Click **Start**. Progress information is displayed in the status line.

   **Note:** If the proprietary communication protocol is used, **Transferring Data** is displayed on the connected Hemochron Signature Elite instrument during data transfer. If the connection is not made, an error message is displayed on the personal computer. Select **Retry** on the error message box to make the connection.

   **Note:** If the POCT-1A communication protocol is used, specific messages indicate the progress of data transfer. If the connection is not made, the instrument will beep, display **Network Timed Out!**, and prompt that the instrument is ready to accept a cuvette.

8. A message box is displayed when writing of the configuration data is complete. Click **OK** to acknowledge the message.

**Printing Instrument Configuration Data**

All of the configuration parameters for an individual instrument or a configuration file can be printed.

1. (If needed) Connect the instrument to the PC (see page 46).
2. (If needed) Start Hemochron **Configuration Manager** and select the configuration file that is to be downloaded (see page 49) or click on **Read Configuration From Instrument**.

---

64
3. Click the **View/Print Configuration** button on the Configuration Module dialog box (page 51) to display the configuration report.

4. Click the Print icon 📜 to print the report.
   - Click the Export icon 📀 to export the report as a specified file type to a specified drive.
   - Click the Refresh icon 🔄 to update the report.

5. Click the Cancel icon ✗ to close the report window.

**Closing the Configuration Module**

After configuration data has been processed, the configuration module can be closed.

1. Click the **Exit** button on the Configuration Module dialog box (page 51) to cancel display of the configuration module dialog box.

**Displaying and Printing Specific Configuration Reports**

Several types of configuration reports can be displayed and printed. The reports provide an audit trail for specific information from all configuration files (or for all instruments) stored in the database:

<table>
<thead>
<tr>
<th>Specific Report</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Instrument Options Listing</td>
<td>Lists instrument options.</td>
</tr>
<tr>
<td>Operator Table Listing</td>
<td>Lists the name, OID, PIN, and certification expiration date for each operator.</td>
</tr>
<tr>
<td>QC Lock-Out Control Listing</td>
<td>Lists selections for EQC/LQC Lockout.</td>
</tr>
<tr>
<td>User Notes Listing</td>
<td>Lists user notes that have been defined.</td>
</tr>
<tr>
<td>Lot Number Listing (Signature Elite v2.0 or higher only)</td>
<td>Lists lot numbers and expiration dates for cuvettes that have been defined for each assay. Lists lot numbers, expiration dates, and expected ranges for controls that have been defined for each assay.</td>
</tr>
</tbody>
</table>

**Audit Trail Report**

Lists the changes made to configuration data files. The report can be sorted by date or user name.

**To Prepare a Specific Configuration Report:**

1. Select **Reports** in the Configuration Manager main menu.
2. Select the desired report type from the **Reports** menu. A drop-down menu is displayed for selecting the instrument type (*Signature Elite* or *Signature+*).
3. Select the instrument type. The specified information is displayed.
4. Click the Print icon 📜 to print the report.
   - Click the Export icon 📀 to export the report as a specified file type to a specified drive.
   - Click the Refresh icon 🔄 to update the report.
5. Click the Cancel icon ✗ to close the report window.
MASTER FILE MAINTENANCE
The Master File Maintenance function of Hemochron Configuration Manager allows a record for a Hemochron Signature Elite instrument or IP address to be added, modified, or deleted.

Manually Adding an Instrument
The supervisor can add a new instrument to the table of instrument serial numbers and unit locations.

1. Select **Serial Number Master File** from the **Master File Maintenance** menu. The Serial Number Master File Maintenance dialog box is displayed:

![Serial Number Master File Maintenance dialog box](image)

2. Click **Add** to display a blank information form.
3. Enter the serial number, unit location, and comment, if desired. The Date Manufactured is read from the instrument.
   
   **Note:** Entries are not case-sensitive.

4. Click **Save**. The new instrument record is saved.

Automatically Adding an Instrument
The Serial Number Master File is automatically updated whenever configuration data is read from an instrument that is not currently included in the Serial Number Master File. A message box is displayed to confirm the update:

![Automatic update message box](image)

Click **OK** to acknowledge the update.
Updating an Instrument Record
The supervisor can modify or delete a record for an existing instrument.

1. Select **Serial Number** from the **Master File Maintenance** menu. The Serial Number Master File Maintenance dialog box is displayed listing information for the first instrument record in the database.
   
   **Note:** The record number and total number of records in the database are displayed at the bottom of the dialog box.

2. Use the scroll buttons at the bottom of the dialog box to display the record to be updated.
3. Update the information as needed. Click **Delete** to delete the instrument record.
   
   **Note:** Entries are not case-sensitive. The Serial Number field and Date Manufactured field cannot be changed. To change the serial number, delete the record and add a new record for the new serial number.

4. Click **Save**. The updates are saved.
   
   **Note:** Click **Exit** to return to the main window.

**CONFIGURATION MANAGER UTILITIES**
The Hemochron Configuration Manager Utilities allow supervisors to maintain the HCM database:

<table>
<thead>
<tr>
<th>Function</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>System Configuration</strong></td>
<td>Update institution name, address, and contact information. Enter the PC COM port used, or automatically detect it.</td>
</tr>
<tr>
<td><strong>Security</strong></td>
<td>Update or delete HCM user records, including user name file, password, or security level.</td>
</tr>
<tr>
<td><strong>Set Instrument Date and Time</strong></td>
<td>Synchronize date and time between an instrument and the PC.</td>
</tr>
<tr>
<td><strong>Purge Configuration Data</strong></td>
<td>Remove a configuration file from the database.</td>
</tr>
<tr>
<td><strong>Rebuild Database</strong></td>
<td>Compress and repair the specified tables. (Compress reduces the size of database files. Repair re-indexes each table).</td>
</tr>
<tr>
<td><strong>Convert HCM Data from Ver 1.0 or Ver 2.0</strong></td>
<td>Converts data from Hemochron Configuration Manager V1.0 or V2.0 to V3.0.</td>
</tr>
</tbody>
</table>

Updating Laboratory Information and/or PC COM Port

1. Select **System Configuration** from the Utilities menu. The System Configuration dialog box is displayed:

   ![System Configuration Dialog Box]

   **Note:** Refer to page 47 for instructions on entering information and selecting a COM port.
Adding a User
The supervisor can add a new user to the table of operators of Hemochron Configuration Manager. The new user will be assigned a password and access level.

Note: Any user can perform all operations until at least one user with an access level of Supervisor is defined. Once a supervisor is defined, only the supervisor can create and edit any files in Configuration Manager. All other users will only be allowed to view and print.

1. Select Security from the Utilities menu. The User Master File Maintenance dialog box is displayed:

2. Click Add to display a blank user information form.
3. Enter the user name, password, e-mail address and other information.

   Note: Entries are not case-sensitive.
4. Select Yes in the Supervisor field if the user is to have supervisor-level access.
5. Click Save. The new user record is saved.

Updating a User Record
The supervisor can modify or delete a record for an existing user.

1. Select Security from the Utilities menu. The User Master File Maintenance dialog box is displayed listing information for the first user record in the database.

   Note: The record number and total number of records in the database are displayed at the bottom of the dialog box.
2. Use the scroll buttons at the bottom of the dialog box to display the record to be updated.
3. Update the name, password or access level as needed. Click Delete to delete the user record.

   Note: Entries are not case-sensitive. The User Name field cannot be changed. To change User Name, delete the record and add a new record for the new User name.
4. Click Save. The updates are saved.

   Note: Click Exit to return to the main window.

Important: The user name “Supervisor” cannot be deleted from the database; however the password can be changed.
Setting the Instrument Date and Time

The time clock of the Hemochron Signature Elite instrument must be synchronized with that of the Hemochron Configuration Manager PC so that correct starting times are used for system lockouts.

Note: The time and date format that are used on an instrument may not be the same as that used in Hemochron Configuration Manager.

Setting the Instrument Date and Time when the Communication Protocol is Used:
1. Start Hemochron Configuration Manager (see page 47).
2. Select Set Instrument Date and Time from the Utilities menu. The Read Instrument Date and Time dialog box is displayed:

   ![Read Instrument Date & Time dialog box]

3. Designate the COM port by selecting it from the drop-down list.
   Note: The instrument must be connected to a COM port. If the instrument is connected to a network, the date and time must be set at the instrument as described on page 19.
4. Click Start. Progress information is displayed in the status line.
   Note: Transferring Data is displayed on the connected Hemochron Signature Elite instrument during data transfer.
   Note: If the connection is not made, an error message is displayed on the personal computer. Select Retry on the error message box to make the connection.
5. At the completion of data transfer, the Set Instrument Date and Time dialog box is displayed:

   ![Set Instrument Date & Time dialog box]

   Note: The Instrument information is read-only and cannot be changed by the user.
6. Set the date and time using the up and down arrow keys. Or, click Use PC Date and Time to display the PC date and time.
7. Click Send to Instrument to synchronize the connected instrument with the displayed date and time.
Setting the Instrument Date and Time when the POCT-1A Communication Protocol is Used:

1. Start Hemochron Configuration Manager (see page 47).
2. Read the configuration from the instrument that is connected over the network (see page 49).

   **Note:** When the POCT-1A communication protocol is used, Hemochron Configuration Manager cannot communicate with the instrument over the COM port.

3. Click the **Clone Instrument** button on the Configuration Module dialog box (page 51) to display the Clear Instrument Counts dialog box:

4. Select **Synchronize Instrument Date + Time with PC (POCT-1A)**. Do not select the other two options.
5. Select **OK**. The Write Configuration dialog box is displayed (page 63).
6. Click **Start**. Progress information is displayed in the status line.
7. A message box is displayed when writing of the configuration data is complete. Click **OK** to acknowledge the message.

**Purging Configuration Data**

An existing configuration file can be removed from the Configuration Manager database. A record of the file removal is maintained in the audit trail.

1. Select **Purge Configuration Data** from the **Utilities** menu. The Purge Configuration Data dialog box is displayed:

2. Use the drop-down arrow to select the configuration file that is to be purged. The information for the configuration file is displayed.

   **Note:** The information is read-only and cannot be changed by the user.

3. Click **OK**. The selected configuration file is purged from the database.

**Rebuilding the Database**

1. Select **Rebuild Database** from the **Utilities** menu. The Rebuild Database dialog box is displayed:
2. Select the operation (Compress Database, or Repair and Compress Database) that is to be performed.
3. Click OK. The databases are repaired and/or compressed.

Converting the Database
1. Select Convert HCM Data from Ver 1.0 (or 2.0) from the Utilities menu. The Convert HCM Data from Version 1.0 (or 2.0) dialog box is displayed:

   ![Convert HCM Data from Version 2.0](image)

   Note: The source and destination paths are shown. These cannot be modified.
2. Click OK to start the conversion. A progress bar is displayed during the conversion, and a message box confirms the data is successfully converted.
### Configuration Manager Error Messages

<table>
<thead>
<tr>
<th>Problem</th>
<th>Cause</th>
<th>Corrective Action</th>
</tr>
</thead>
<tbody>
<tr>
<td>CHKSUM error during data transfer</td>
<td>Other software applications are open during data transfer.</td>
<td>Close all applications including anti-virus programs during communications.</td>
</tr>
<tr>
<td>Communication error/ Initialization error</td>
<td>Incorrect cable is used or cable is not completely connected.</td>
<td>Ensure proper cable is being used. Check cable connections.</td>
</tr>
<tr>
<td>No COM Port</td>
<td>No COM Port available.</td>
<td>Use HCM to detect available COM Port</td>
</tr>
<tr>
<td>Text crowded / Overlapping on screen</td>
<td>Older computer graphics card</td>
<td>Does not affect program. Printouts will be correct.</td>
</tr>
</tbody>
</table>

### Instrument Error Messages

The error messages that may be displayed while operating the Hemochron *Signature Elite* instrument are listed below. The probable cause and corrective action are shown for each message. Some messages designate a test or sample fault. This message is included in the printed results for the affected sample. A list of these messages can be found in *Acronyms for Error Messages* on page 43.

If needed, contact Technical Support by phone at 1-877-579-4441 (toll free), 1-858-263-2502 (International), or by FAX at +1-858-314-6700, or by e-mail at techsupport@accriva.com.

<table>
<thead>
<tr>
<th>Error Message</th>
<th>Cause</th>
<th>Corrective Action</th>
</tr>
</thead>
<tbody>
<tr>
<td>RTC....FAULT</td>
<td>Can not communicate with external Real Time Clock.</td>
<td>Contact Technical Support.</td>
</tr>
<tr>
<td>CHARGE BATTERY</td>
<td>Battery power depleted below predetermined level.</td>
<td>Connect to AC/DC Power Module and charge battery for 8 hours.</td>
</tr>
<tr>
<td>BATTERY FAULT</td>
<td>Battery is discharged.</td>
<td>Charge battery for 8 hours. If message persists, contact Technical Support.</td>
</tr>
<tr>
<td>BATTERY TOO HIGH</td>
<td>Battery voltage exceeds 8.8 volts.</td>
<td>Disconnect the AC/DC Power Module. Contact Technical Support.</td>
</tr>
<tr>
<td>Heater Too Cool</td>
<td>Incubator remained below 36 °C after 90 seconds of warming on external power or for up to 150 seconds on battery.</td>
<td>Repeat test, then charge battery. If message persists, contact Technical Support.</td>
</tr>
<tr>
<td>Heater Too Hot</td>
<td>Incubator exceeded 38 °C for 2.5 seconds.</td>
<td>Repeat test with new cuvette. If message persists, contact Technical Support.</td>
</tr>
<tr>
<td>Error Message</td>
<td>Cause</td>
<td>Corrective Action</td>
</tr>
<tr>
<td>-----------------------</td>
<td>------------------------------------------------------</td>
<td>-----------------------------------------------------------------------------------</td>
</tr>
<tr>
<td>Detector Fault</td>
<td>Light path between LED and detectors is blocked.</td>
<td>Repeat test with new cuvette. If message persists, contact Technical Support.</td>
</tr>
<tr>
<td>Detector Blocked</td>
<td>EQC is being run while a cuvette is inserted.</td>
<td>Remove the cuvette and repeat the EQC.</td>
</tr>
<tr>
<td>Sample Pos Fault</td>
<td>Sample has moved outside of testing area in cuvette.</td>
<td>Contact Technical Support.</td>
</tr>
<tr>
<td>ASSAY LOCKED CALL SUPERVISOR</td>
<td>The QC failure limit has been exceeded.</td>
<td>Remove the cuvette. Contact the supervisor.</td>
</tr>
<tr>
<td>MACHINE LOCKED CALL SUPERVISOR</td>
<td>The EQC failure limit has been exceeded.</td>
<td>Contact the Supervisor.</td>
</tr>
<tr>
<td>Sample Not Seen</td>
<td>Sample has not reached front detector in specified time period.</td>
<td>Repeat test with new cuvette. If message persists, contact Technical Support.</td>
</tr>
<tr>
<td>Cuvette Removed</td>
<td>Cuvette was prematurely removed from instrument while testing in progress.</td>
<td>Repeat test with new cuvette.</td>
</tr>
<tr>
<td>Sample Too Large/Sample Too Small</td>
<td>Excess or insufficient sample.</td>
<td>Repeat test with new cuvette.</td>
</tr>
<tr>
<td>Premature Sample</td>
<td>Sample was detected at front detector before specified time period. May occur if sample is added before pump-priming sequence is complete.</td>
<td>Repeat test with new cuvette. If message persists, contact Technical Support.</td>
</tr>
<tr>
<td>MEMORY FAULT</td>
<td>Malfunction in the computer's memory.</td>
<td>Contact Technical Support.</td>
</tr>
<tr>
<td>START Timed Out</td>
<td>START key was not pressed within 5 minutes after entering ready mode.</td>
<td>Remove cuvette and repeat test with new cuvette.</td>
</tr>
<tr>
<td>Unsupported Assay</td>
<td>Cuvette cannot be identified by instrument.</td>
<td>Remove cuvette and repeat test with new cuvette. Use Hemochron cuvettes only.</td>
</tr>
<tr>
<td>No Data Stored</td>
<td>Operator attempted to print data when none is stored in database.</td>
<td>N/A</td>
</tr>
<tr>
<td>User Abort</td>
<td>The test was aborted by the user.</td>
<td>Repeat test.</td>
</tr>
<tr>
<td>Invalid Lot #</td>
<td>The barcoded cuvette or QC lot number label that was scanned was not recognized.</td>
<td>Check the label for damage, then repeat the scan.</td>
</tr>
<tr>
<td></td>
<td>The incorrect format was entered for the lot number.</td>
<td>Repeat the entry using the correct format.</td>
</tr>
<tr>
<td></td>
<td>The lot that was scanned does not match the cuvette test type.</td>
<td>Verify the test type, then rescan the lot.</td>
</tr>
<tr>
<td>Error Message</td>
<td>Cause</td>
<td>Corrective Action</td>
</tr>
<tr>
<td>---------------------</td>
<td>------------------------------------------------------------------------</td>
<td>------------------------------------------------------------------------------------</td>
</tr>
<tr>
<td>Lot Expired</td>
<td>The cuvette and/or QC material that is being used has reached its expiration date.</td>
<td>Remove the cuvette and repeat the test using supplies that are within their expiration date.</td>
</tr>
<tr>
<td>Action Denied</td>
<td>Date/Time cannot be changed if QC Lockout is required, or Date/Time function is denied through Configuration Manager.</td>
<td>Supervisor needs to reconfigure instrument using HCM.</td>
</tr>
<tr>
<td>Disallowed Assay</td>
<td>Assay performance has been prohibited via Configuration Manager.</td>
<td>Supervisor needs to reconfigure instrument using HCM.</td>
</tr>
<tr>
<td></td>
<td>The table of cuvette or QC lots is empty or contains only expired lots.</td>
<td>Supervisor needs to reconfigure instrument using HCM.</td>
</tr>
<tr>
<td></td>
<td>The QC interval is set to zero in Configuration Manager.</td>
<td>Supervisor needs to reconfigure instrument using HCM.</td>
</tr>
<tr>
<td>Cannot erase DB</td>
<td>Erasing patient and/or QC database is not allowed.</td>
<td>Supervisor needs to reconfigure instrument using HCM.</td>
</tr>
<tr>
<td>INR&lt; 0.8</td>
<td>Test result is outside clinical range.</td>
<td>Repeat test with new cuvette.</td>
</tr>
<tr>
<td>INR &gt; 10.0</td>
<td>Test result is outside clinical range.</td>
<td>Repeat test with new cuvette.</td>
</tr>
<tr>
<td>APTT &lt; 20</td>
<td>Test result is outside clinical range.</td>
<td>Repeat test with new cuvette.</td>
</tr>
<tr>
<td>Out of Range-Lo</td>
<td>Sample has clotted prematurely, or did not mix correctly in cuvette.</td>
<td>Repeat test with new cuvette.</td>
</tr>
<tr>
<td></td>
<td>Bubbles may be present.</td>
<td>Repeat test with new cuvette.</td>
</tr>
<tr>
<td>Out of Range-Hi</td>
<td>Test result is outside clinical range.</td>
<td>Repeat test with new cuvette.</td>
</tr>
<tr>
<td>Check Time/Date</td>
<td>Low Battery or RTC has lost its Time/Date Tracking.</td>
<td>Enter or verify Time &amp; Date. Charge Battery.</td>
</tr>
<tr>
<td>Dark Photo Fault</td>
<td>Hardware malfunction.</td>
<td>Contact Technical Support.</td>
</tr>
<tr>
<td>Unsupported POCT &gt;&gt; COM</td>
<td>The POCT-1A communication protocol was attempted to be set while the COM Port is selected, or vice versa.</td>
<td>Set POCT-1A to run over NET, or set to run over COM or NET.</td>
</tr>
</tbody>
</table>
Verification of Barcode Scanner

Check the barcode scanner by scanning a test barcode.

To Test the Barcode Scanner:

1. Press PRINT/SCAN to display the Print menu:

   ![Print Menu 1]

2. Press 7 to display the second page of the Print menu:

   ![Print Menu 2]

3. Press 3 to test the scanner:

   ![Test SCAN]

4. Position the scanner port (see page 5) approximately four inches from the barcode label that is shown below:

   ![Barcode Image]

   **Note:** The barcode shown above is formatted as Code 128.

5. Press PRINT/SCAN. The characters that are read from the barcode are displayed in the screen:

   ![Test SCAN]

6. Verify that **test1234567890abcdefgxyztest** is displayed.

7. Press CANCEL three times.
SERVICE AND MAINTENANCE

Routine Maintenance
Inspect and clean the cuvette opening as required. Remove residual dried blood or other foreign matter using water-moistened cotton swabs.

Remove any residual water with dry cotton swabs. If a disinfectant is needed, use a 0.5% solution of sodium hypochlorite or a 10% dilution of household bleach in water. Wipe instrument with a water-dampened cloth to remove residual bleach from the plastic surfaces.

Apply solution to clean and disinfect areas contaminated with blood. DO NOT use solvents or strong cleaning solutions as they may damage the instrument’s plastic components. Routine maintenance other than cleaning normally is not required.

It is recommended to clean and decontaminate prior to service or transfer of equipment.

Service
The Hemochron Signature Elite instrument is almost completely self-monitoring and has no user serviceable parts.

It monitors internal circuitry and reports problems to the display screen automatically. Malfunctions are indicated by error messages detailed in the section “Troubleshooting.”

Battery Care
To optimize battery life, it is recommended that the Hemochron Signature Elite instrument be run on its battery during the day. It can be plugged in overnight to allow the batteries to recharge. The life of the lithium battery is optimized when the battery is exercised in this manner.

A fully charged battery will operate as defined in the Operation specifications section on page 12. When the batteries are drained to the point that valid testing cannot be performed, the instrument will display “CHARGE BATTERY” (see “Troubleshooting”). At this point, the instrument must be plugged in for operation and recharging. Once plugged into a Hospital Grade AC outlet, the instrument can be used immediately.

Caution
The battery used in this device may present a risk of fire or chemical burn if mistreated. Do not disassemble, heat above (60°C), or incinerate. Use of another battery may present a risk of fire or explosion.

Dispose of used battery promptly. Keep away from children. Do not disassemble and do not dispose of in fire.

Instrument Disposal
If disposal is necessary, follow local regulations for disposal of Li-Ion batteries and electronic devices.
SAFETY STANDARDS

The Hemochron Signature Elite instrument complies with the following safety standard requirements and directives:

CAN/CSA C22.2 No. 61010-1:2012
CAN/CSA C22.2 No. 61010-2-101:2009
CAN/CSA C22.2 No. 61010-2-010:2009

UL 61010-1:2012

IEC/EN 61010-1:2010
IEC/EN 61010-2-101:2002
IEC/EN 61010-2-010:2003

IEC 60825-1:2007

EN 55011:2009 A1:2010 Group 1 Class A
EN 61326-1:2006 Class A limit

This instrument has been designed and tested to CISPR 11 Class A. In a domestic environment it may cause radio interference, in which case, you may need to take measures to mitigate the interference.

It is the manufacturer’s responsibility to provide electromagnetic compatibility information to the customer or users. It is the user’s responsibility to insure that a compatible electromagnetic environment for the instrument can be maintained in the order that the device will perform as intended.

EN 61326-1:2006 Table 2 limits

Do not use this instrument in close proximity to sources of strong electromagnetic radiation (e.g. unshielded intentional RF sources), as these may interfere with proper operation.

Protection Against Ingress of Liquids: Ordinary (no protection as defined by IEC 60529)
Product Cleaning and Disinfection: Only according to recommendations of the manufacturer’s accompanying documentation
Mode of Operation Of Equipment: Continuous
Degree of Safety of Application in the Presence of Flammable Anesthetic Mixture With Air, Oxygen or Nitrous Oxide: Not Suitable

NOTE: as defined in the above standards, the classification of Not Suitable is not intended to indicate that the instrument is not suitable for use in an Operating Room (OR) environment. Rather, it is intended to indicate that the instrument is not suitable for use in the direct presence of a flammable anesthetic mixture with air, oxygen or nitrous oxide.

All relevant documentation is kept on file at Accriva Diagnostics, Inc., San Diego, CA.
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INDEX

acronyms used .......................................... 4
action keys.................................................. 5
adding a user ............................................. 72
attention label ............................................. 11
battery
  care.......................................................... 80
  charging................................................... 15
  checking................................................... 16
  life.......................................................... 13
  low warning............................................. 16
  type........................................................ 13
baud rate................................................... 21
cable configurations............................... 27
charging battery....................................... 15
CLSI Standard POCT-1A................................. 21
commands.................................................. 8
communication parameters............... 22, 47
computer requirements ................................. 49
configuration
  911 limit.................................................. 61
cuvette lots ............................................... 55, 56
date format ............................................... 55, 57
EQC failure limit ........................................ 61
instrument.................................................. 55
LQC failure limit ........................................ 61
Operator ID and PIN ..................................... 56
operator ID options ................................. 55, 57
patient ID options ................................. 55, 57
preparing the instrument ......................... 49
QC exceptions............................................ 55
QC lockout................................................ 55, 59
QC lots...................................................... 55, 56
requiring cuvette lot entry ....................... 57, 65
requiring QC lot entry................................. 58
specifying EQC level 2 ............................... 60
supervisor PIN ........................................... 56
time format............................................... 55, 57
user notes................................................ 56, 64
configuration options............................... 55
connecting
  computer ............................................... 26
  network.................................................. 26
  printer..................................................... 26
custom header.......................................... 43
cuvette..................................................... 9
cuvette lot number
  entering.................................................. 30, 65
cuvette validation...................................... 38
definitions............................................... 4
deleting a user record................................. 71, 72
description

features................................................... 9
overview.................................................. 5
disallowed assay....................................... 60
display panel............................................. 6
EQC .......................................................... 59
erasing
  patient database.................................... 44
  QC database............................................ 44
error messages......................................... 46, 76
features................................................... 9
Hemochron Configuration Manager
  commands................................................. 52
  installing............................................... 49
  menus.................................................... 52
  starting................................................ 50
ID
  entering operator ID (OID) ......................... 30
  entering patient ID (PID) ........................... 29
instrument options..................................... 56
instrument self-check .................................. 37
instrument settings
  display brightness..................................... 18
  display contrast....................................... 18
  flashlight time........................................ 18
intended use............................................. 2
interface cable.......................................... 26
keypad ..................................................... 5
limitations................................................. 40
liquid controls.......................................... 38
lockout
  instrument............................................ 10
  QC parameters......................................... 59
lot numbers
  entering cuvette lot number ........................ 30
  entering QC lot number............................... 34
maintenance............................................. 80
mandatory entry of
  cuvette lot number.................................. 30, 34
  operator ID (OID)..................................... 32
  operator PIN ........................................... 33
  patient ID (PID) ....................................... 32
materials
  optional................................................ 14
  provided............................................... 14
  required................................................ 14
menus....................................................... 7, 52
note
  addition................................................. 35
  operator ID (OID)
    entering............................................... 30
    entering a mandatory operator ID... 32
overview ............................................... 2
patient ID (PID) ........................................ 20
    entering ........................................... 29
    entering a mandatory patient ID ....... 32
prewarm .............................................. 10
printing .............................................. 40
    all patient results for a cuvette lot .. 41
    all patient results for an assay ...... 41
    all patient results from a specific date 41
    all patient results from a specific operator ....... 41
    all QC results for a cuvette lot .... 42
    all QC results for an assay ........ 42
    all QC results from a specific date 42
    all results for a specific patient .... 40
    all results in a database ............. 40
    QC all QC results from a specific operator ................................................................................ 41
    QC results for a QC lot .............. 42
    results of last test ...................... 40
system information ......................... 43
QC lot number ........................................ 20
    entering ........................................... 34, 66
Quality Control ...................................... 37
    electronic (EQC) ................................ 37
    instrument performance ................ 37
    instrument temperature ................ 37
    liquid (LQC).................................... 38
    liquid controls .............................. 37
results management .............................. 40
printing .............................................. 40
reviewing .............................................. 40
    patient database .............................. 44
    QC database ..................................... 45
running a test ....................................... 28
scanning...................................................... 40
Operator ID (OID) .................................. 30
Operator IDs ........................................ 20
patient ID (PID) ........................................ 20
QC lot numbers ........................................ 20
service .................................................. 80
setting ..................................................... 17
    date .................................................. 17
    time ................................................. 16
shutdown ............................................... 17
specifications .......................................... 37
    AC/DC Power Module ....................... 13
    dimensions ...................................... 13
    performance ..................................... 13
    weight ............................................. 13
    specifying QC lockout ..................... 59
    specimen collection ......................... 36
starting a test ......................................... 36
starting the instrument ...................... 27
supervisor .............................................. 37
    functions ......................................... 19
    PIN .................................................. 55, 56
terminology .......................................... 40
test ...................................................... 40
    running ........................................... 28
    starting ........................................... 36
test cuvettes ......................................... 36
    test timeout ..................................... 10
time and date ........................................ 16
time zone .............................................. 21
troubleshooting ..................................... 76
updating a user record ........................ 71
    test cuvettes ..................................... 9
    test timeout ..................................... 10
    time and date ................................... 16
    time zone .......................................... 21
    troubleshooting .................................. 76
    updating a user record .................... 71
    updating laboratory information ....... 71
    updating PC COM port ....................... 71
user ..................................................... 72
    deleting ........................................... 72
    updating ......................................... 72

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