VITROS® ECiQ Immunodiagnostic System

- Featuring Intellicheck® Technology for the highest level of confidence and security in patient results
- Simple to use, with continuous, STAT, and random access testing
- Proprietary Enhanced Chemiluminescence detection technology provides wide dynamic ranges and exceptional precision and sensitivity







VITROS® ECiQ Immunodiagnostic System Specifications

Measurement Principle: Enhanced Chemiluminescence

MicroWell Technology: Excellent assay sensitivity and precision, small sample volumes and minimal waste

Reagents: No preparation; no mixing and reconstitution required for Integrated Reagent Packs, Signal Reagent and Universal Wash Reagent

On-Board Test Capacity:

- Up to 2000 assays
- Up to 20 different Integrated Reagent Packs
- 100 assays each
- · Resident on-board simultaneously
- Automatic, self-contained, temperature and humidity controlled Reagent Supply

Calibration: Multiple lots may be pre-calibrated with automatic lot switching

- Up to 28 days
- Process: Random Access Calibration with Automatic Result Protection; 25 calibrations across 16 lots per assay

System Startup: Automatic Integrated Prime/Purge

- No primes, purges, washes or tubing maintenance
- No daily calibrations or calibration checks

Throughput: Up to 90 reportable patient results per hour

Sample Types: Serum, Plasma, Urine, Amniotic Fluid*, Whole Blood**

Sample Volume: 10-80µL

Sample Capacity: 60 samples in Universal Sample Trays

Sample and Reagent Management:

Intellicheck® Technology

- Disposable Tip Metering verifies sample aspiration and dispense and addresses carryover and cross-contamination concerns
- Clot, bubble, low and high viscosity, thin layer fluid and short sample
- $\bullet \ \, Save\text{-the-Sample}{}^{TM} \ \, Clot/Bubble \\$ Management
- · Liquid level sensing
- MicroWell Dispense Verification
- Reagent Aspiration and Dispense Verifications

Sample Containers: Universal Sample Trays accommodate:

- 10mL, 7mL, 5mL collection tubes
- 1.5mL micro-collection containers
- VITROS Microsample cups and 0.5mL and 2.0mL cups

Sample Bar Code Identification:

Autodiscriminates by simultaneously recognizing all standard symbologies:

- Code 128
- ISBT 128
- Code 39
- Codabar
- Interleaved 2 of 5

Automatic Dilution:

- Reflex dilution
- Operator requested dilution
- · Protocol and pre-treatment dilutions

Automatic Reflex Testing:

- Reflex to different assays
- · Reflex to the same assay

Automatic Repeat Testing: Repeats samples automatically after a result is not initially reported for the sample

Operator Interface: Color-coded graphical user interface

- · Ergonomic flat, low-glare, LCD, touchscreen monitor
- Keyboard platform and support arm
- Flexible, customized positioning
- On-board documentation and Help

System Dimensions:

Width: 111.8 cm / 44 inches Depth: 73.7 cm / 29 inches Height: 130.2 cm / 51.25 inches Weight: 366 kg / 807 pounds

POWER

Line Voltage: Dedicated, single phase AC power line North America: 120 V AC

Continental Europe: 200-240 V AC

Line Frequency:

North America: 50-60 Hz Continental Europe: 50-60 Hz

ENVIRONMENT

Average Heat Output: 4,100 BTUs per hour Operating Temperature: 15°-30°C / 59°-86° F Ambient Relative Humidity: 15%-75% RH noncondensing

Altitude: -0.1524/2.439 km / -500/8000 feet

Plumbing: No water or drain required; Selfcontained, on-board liquid waste management eliminates special requirements for off-board plumbing

COMMUNICATIONS

Bidirectional interface for ASTM and Kermit protocols, including broadcast download

4 RS 232 serial ports

e-Connectivity® Interactive System Management

- · Using a dedicated analog telephone line and a modem, a VPN establishes a secure connection between an enabled ECiQ System and Ortho-Clinical Diagnostics Technical Support
- · Automatic Two-Way Data Exchange to automatically send and retrieve data. Includes automatic download of system software updates.
- · Remote Connectivity provides the ability to enable Remote Diagnostics and Remote Control operation

*Some or all types of specimens or suggested reference interval or cutoff for these analytes are not approved or cleared for market in the United States

For more information contact your Ortho-Clinical Diagnostics representative or visit our website at www.orthoclinical.com.