



CS-2500 System

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Advance with Confidence

The CS-2500 System leverages PSI™ technology with primary-tube sample-volume checks and double aspiration clog detection to identify and automatically manage potentially problematic test samples prior to analysis, minimizing repeat testing. Designed for labs that run fewer than 300 hemostasis samples per day, the CS-2500 System provides lab-to-lab consistency for multisite patient monitoring, with sample result traceability for in-depth audit capabilities and sophisticated cap-piercing technology for dependable throughput under all conditions. With low maintenance, onboard consolidation of a variety of test methods, and flexible reagent management, the CS-2500 System is an affordable and efficient solution for your lab.

Reagent Test Menu

| | |
|-----------------------|--|
| PT | Dade® Innovin®, Thromborel® S |
| APTT | Dade Actin FS®, Dade Actin FSL |
| Fibrinogen | Dade Thrombin |
| Thrombin time | Test Thrombin |
| Batroxobin time | Batroxobin |
| Factor deficiency | Factor II, V, VII, VIII, IX, X, XI, XII |
| Lupus anticoagulant | LA1 screening, LA2 confirmation |
| Protein C pathway | Protein C, F V Leiden, INNOVANCE® Free Protein S |
| Anti-Xa | INNOVANCE Anti-Xa, INNOVANCE Apixaban |
| Antithrombin III | INNOVANCE Antithrombin |
| D-dimer | INNOVANCE D-Dimer |
| Von Willebrand factor | INNOVANCE VWF Ac, INNOVANCE VWF Ag |
| Chromogenic | Berichrom® Factor VIII Chromogenic, Berichrom α2-Antiplasmin, Berichrom Plasminogen, Berichrom Protein C |



CS-2500

System Specifications

Measurement

| | |
|--|---|
| Principle | Photo-optical based on change in the transmitted light emitted from the sample with added reagent |
| Method/channels | Clotting (10 single channels); chromogenic, immunoassay, and aggregation methods [†] |
| Measurement channels | 10 wells (mixing function using stir bars, possible for 4 wells) |
| Source lamp | Halogen lamp for measurements at 340, 405, 575, 660, and 800 nm wavelengths |
| Analysis mode | Normal and microsample modes |
| Analysis method | Dilution analysis; automatic reanalysis (redilution analysis, reanalysis, reflex testing) |
| Time resolution | Sampling can be performed at intervals of 0.1 seconds |
| Measuring time | Up to 1800 seconds for each parameter |
| Incubation | 10 wells |
| Number of user-definable (open) channels | 80,000 |

Sample Handling

| | |
|------------------------------|---|
| Type of sample | Primary tubes and/or sample cups |
| Sampling mechanism | Automated sample and standard predilution |
| Sample integrity check (PSI) | Tube fill vol check, hemolysis and lipemia check by assay |
| Traceability of results | Operator name, test date and time, reagent lot information, test reaction position, testing and reagent table temperatures, test protocol number, dilution ratio, QC performed date, calibration curve identification, and maintenance logs |
| Carrier system | Continuous-access sample racks with 10-tube capacity |
| Maximum load | 50 samples; 5 racks with 10 samples per rack |
| STAT sample loading | Five priority positions |
| Storage temperature | Room temperature |
| Racks various | Five sample racks, continuous loading; tube holders and adapters available |
| Handling | Flexible mix of capped and/or uncapped sample tubes and 4 mL conical sample cups |
| Primary sample probe | Liquid-level sensing, crash protection, clog detection, liquid surface verification |
| Cap piercing | Pierces caps on primary tubes |

Reagent Handling

| | |
|------------------------------|--|
| Reagent recognition | Internal reagent bar-code identification |
| Dispensing mechanism | Two probes: one heated for reagents and one for samples, controls, and calibrators Two catchers |
| Reagents onboard | 40 reagent/control positions and 5 buffer/rinse solution positions |
| Storage temperature | 40 cooled reagent positions onboard at 10°C ±2°C |
| Handling | Flexible mix of reagent positions within the reagent table; various adapters available |
| Mixing position | Up to 10 positions available on the reagent table |
| Reagent inventory management | Tracks number of tests remaining, lot number, onboard stability, vial type, set date and time, and expiration date |

Throughput* (tests/hour, approx.)

| | |
|---------------|--|
| PT | 180 (single-parameter analysis) |
| APTT | 180 (single-parameter analysis) |
| PT/APTT | 180 (simultaneous analysis) |
| PT/APTT/AT/DD | 95 (simultaneous results for normal mode processing) |

Operation

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|--------------------------|--|
| Access mode | Continuous random access (samples, reaction tubes) |
| Calibration | Two 12-point calibration curves with maximum of 5 repeated analyses per point and up to 10 calibration curves; one reagent lot group |
| Calibration curve | 250 user-defined parameters |
| View calibration | Graphical display of calibration curves from up to 10 different reagent lots/parameters |
| Auto calibration/auto QC | User-defined time interval or with new reagent vial |
| Temperature control | Measuring channels 37°C ±0.5°C, incubation area 37°C ±1.0°C, reagent probe 37.5°C ±0.5°C |

Reaction Tubes

| | |
|------------------------|---|
| Type | Single reaction tubes |
| Loading | Automatic continuous access, 500 reaction tubes onboard |
| Reaction tube stirring | Yes |

System Fluid

| | |
|----------------------|---|
| Cleaning and rinsing | Washing solutions onboard |
| System containers | 20 L containers for water (rinse) and waste (optional) Rinse may be direct-filled with house water by optional water controller module Waste may be direct-plumbed to drain |

Computer/Printer

| | |
|--------------------------|---|
| Workstation | PC |
| Display | 24" touchscreen |
| Printer | Graphic printer |
| Input devices | Touchscreen, keyboard, and mouse, 2-D bar-code reader |
| Data storage | About 10,000 samples with a maximum of 60 results per sample |
| Onboard maintenance logs | Scheduled and automatically monitored routine maintenance activities via software |

Software

| | |
|-----------------------|--|
| Anti-virus Protection | McAfee Embedded Control |
| LIS interface | CA-1000/1500 System: ASTM CS-2000i System: ASTM1381-95/1394-95, ASTM1381-02/1394-97 |
| Host connection | Bidirectional RS-232C serial port or via Ethernet-TCP/IP |
| Operating system | Microsoft Windows 10 |

Power Supply

| | |
|-------------------|---|
| Operating voltage | Main unit: 100–240 V Pneumatic unit: 100–117 V/220–240 V |
| Power consumption | Main unit: ≤800 VA Pneumatic unit (processing): ≤280 VA |
| Main frequency | 50–60 Hz |

Environmental Conditions

| | |
|------------------------|--|
| Operating temperature | 15–30°C |
| Environmental humidity | 30–85% (no condensation except on the reagent table) |
| Atmospheric pressure | 70–106 kPa |
| Waste heat | Approx. 4000 BTU/h (1040 Kcal/h) |
| Noise level | 60 dB or below (excluding sudden noise that stops within 5 seconds and alarms) |

Dimensions

| | |
|-----------------------|---|
| Main unit and cabinet | Approx. 775 (W) × 895 (D) × 685 (H) mm 30.6 (W) × 35.2 (D) × 27.0 (H) in. |
| Pneumatic unit | Approx. 280 (W) × 355 (D) × 400 (H) mm 11.1 (W) × 14.1 (D) × 15.7 (H) in. |
| Terminal (PC) | Approx. 338 (W) × 381 (D) × 100 (H) mm 13.3 (W) × 15.0 (D) × 3.95 (H) in. |
| Terminal (monitor) | Approx. 558 (W) × 122.5 (D) × 403.5 (H) mm 21.8 (W) × 4.8 (D) × 15.8 (H) in. |

Weight

| | |
|--------------------|-----------------------------|
| Main unit | Approx. 110 kg 242.5 lb |
| Pneumatic unit | Approx. 17 kg 37.5 lb |
| Terminal (PC) | Approx. 6.84 kg 15.08 lb |
| Terminal (monitor) | Approx. 5.6 kg 12.3 lb |

Quality Control

X-control, Levey-Jennings control
Multi-rule monitoring (Westgard rule)
Maximum of 1200 plots × 750 files can be saved

Compliance

| | |
|------------------------------|--|
| Safety standards (main unit) | IEC 61010-1:2001 (2nd Edition) IEC 61010-2-081:2001 + A1:2003 IEC 61010-2-101:2002 IEC 60825-1:1993 + A1:1997 + A2:2001 EN 61010-1:2001 (2nd Edition) EN 61010-2-081:2002 + A1:2003 EN 61010-2-101:2002 EN 60825-1:1994 + A2:2001 + A1:2002 UL 61010-1:2004 (2nd Edition) CAN/CSA-C22.2 No. 61010-1-2004 (2nd Edition) CAN/CSA-C22.2 No. 61010-2-081-04 (R09) CAN/CSA-C22.2 No. 61010-2-101-04 (R09) CAN/CSA-E60825-1-03 |
| EMC standards | IEC 61326-1:2005 IEC 61326-2-6:2005 CISPR 11:2009/A1:2010 Group 1 Class A IEC 61000-3-2:2006/A1:2009/A2:2009 IEC 61000-3-3:2008 IEC 61326-1:2005 (Table 1) IEC 61326-2-6:2005 IEC 61000-4-2:2008 IEC 61000-4-3:2006/A1:2007/A2:2010 IEC 61000-4-4:2004/A1:2010 IEC 61000-4-5:2005 IEC 61000-4-6:2009 IEC 61000-4-8:2009 IEC 61000-4-11:2004 |

For more information, please contact your representative from Siemens Healthineers or visit our website.

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†Aggregation methods under development.

‡Throughput values were determined by the time to first result; processing capability varies depending on the reagent used. Throughput values stated above were determined using Siemens' study protocol with PT (Thromborel S Reagent), APTT (Pathromtin® SL Reagent), INNOVANCE D-Dimer Reagent, and AT (INNOVANCE AT Reagent) test applications.

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