

## **Confidence in Results**

With Data Integrity







## Confidence in Test Results With Data Integrity

Your clinical laboratory must quickly and cost-effectively deliver accurate patient test results to the healthcare professionals you serve. With a simplified user interface and three-tiered access to software with permission levels for technicians, supervisors and directors, Thermo Scientific $^{\text{TM}}$  ClinQuan  $MD^{\text{TM}}$  software for in vitro diagnostic use is designed to help you meet this challenge confidently day after day.

Using ClinQuan MD software, technical staff can store, retrieve and process LC-MS data with ease. Audit trails help ensure the integrity of results for the numerous laboratory-developed LC-MS tests you run.

Combine ClinQuan MD software with the Thermo Scientific<sup>™</sup> Prelude MD<sup>™</sup> HPLC and Endura MD<sup>™</sup> mass spectrometer, and your laboratory can obtain the quantitative accuracy of LC-MS, easily and confidently.

# Store, Retrieve, and Process Data With Confidence

#### Streamline Workflows

ClinQuan MD software features a simplified user interface designed to make routine quantitation simple, fast, and productive. With a step-by-step process for method selection, building sample lists, running samples, and performing daily maintenance, quality control, and reporting, you can be sure users are able to obtain accurate results.

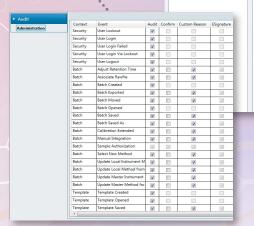




LAB DIRECTOR LOGIN
ADMINISTRATION
CONSOLE

**ASSIGN ROLE** 

**AUDIT** 



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Acquisition Wizard

Allow Sample Reinjection

Extended Calibration

Require Acquisition Checkl

Template Editing

Administrator
Auditing

Results

Data Review

Edit

Local Instrument Method - Update From Master Method
 Local Instrument Method - Update To Master Method

## Address CLIA-Required Roles and Responsibilities

The software features address Clinical Laboratory Improvements Amendments of 1988 (CLIA) with the three CLIA-required permission levels: lab director, supervisor (technical consultant), and technician (testing personnel). The lab director maintains the flexibility to tailor task permissions to specific staff according to their qualifications, training, skills, or to specific laboratory standard operating procedures.



#### **Process Test Results**

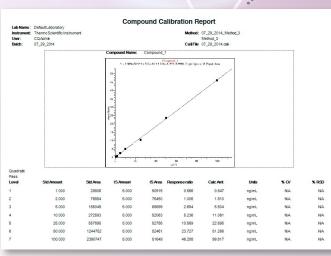
SUPERVISOR LOGIN

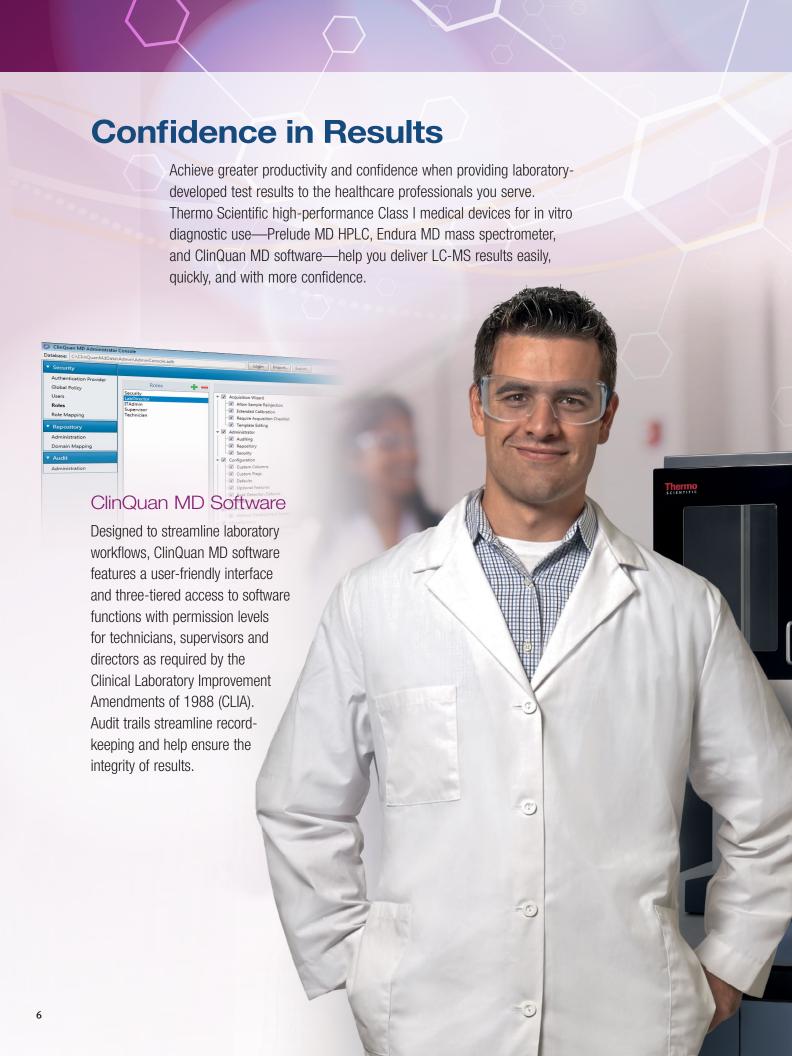
REVIEW TEST RESULTS

APPROVE AND FILE REPORT

### Maintain Records, Help Ensure Data Integrity

In order to streamline record keeping and ensure the integrity of results, records are reviewed, approved, and stored with a time-stamped audit trail. ClinQuan MD software records all user actions, from method building to reviewing and reporting data.









For in vitro diagnostic use. Not available in all countries. Please visit www.thermoscientific.com/clinquanMD For complete product line information visit www.thermoscientific.com

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