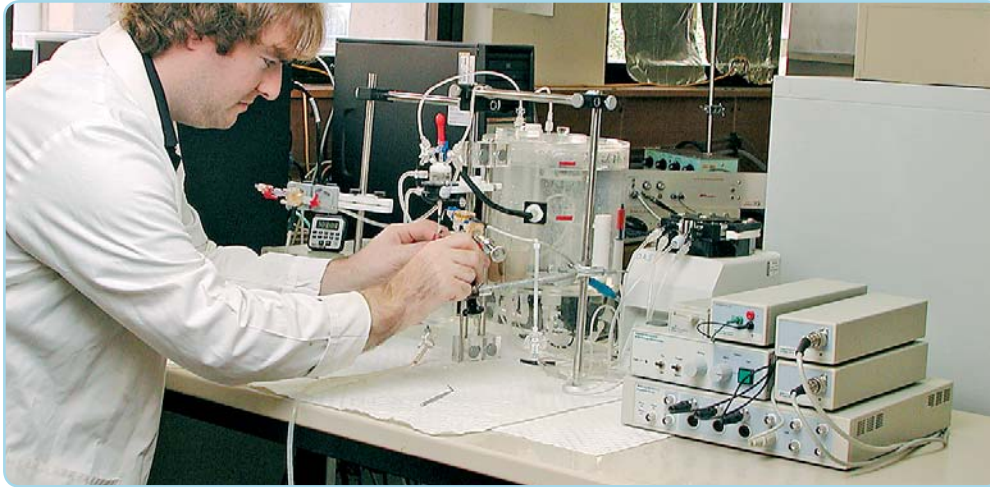


GLP & 21 CFR Part 11 Compliance

Data Acquisition Solutions



Meeting the requirements of GLP does not need to be a time-consuming or frustrating exercise. ADInstruments offers an easy and reliable data acquisition solution for a GLP and 21 CFR Part 11 compliant environment.

The ADInstruments GLP Client enables you to satisfy the requirements for non-repudiation of data. Features include a comprehensive audit trail, secure signing, preservation of raw data and date/time stamping. The GLP Server software provides a centralized user authorization system. The two GLP software components work with any PowerLab® data acquisition unit and LabChart® software to facilitate GLP and 21 CFR Part 11 compliance.

Installation, Validation & Training

ADInstruments offers installation and training services that incorporate the use of hardware and software to optimize data acquisition and analysis. We also provide services for periodic calibration of your PowerLab hardware to confirm that the system is operating within manufacturing specifications. The ADInstruments Validation Unit is available for customers wishing to perform in-house validation of PowerLab systems. It includes all the hardware, software programs and documentation required for validation purposes.

Online Support & Hardware Warranty

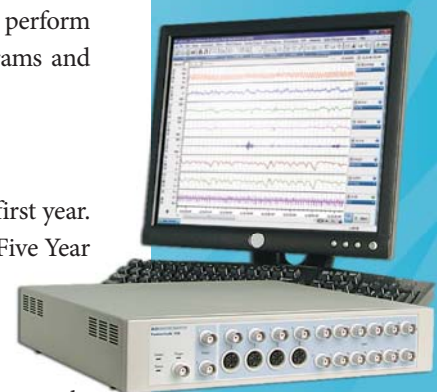
Free email, phone and online GLP technical support is available to all customers in the first year. All PowerLab data acquisition systems come with a standard three year warranty. A Five Year Extended Warranty and Support Agreement is also available. Contact us for details.

Proven Quality & Reliability

ADInstruments is a leader in computer-based data acquisition systems for life science research. We have installed over 23000 PowerLab systems worldwide in pharmaceutical companies, research organizations, hospitals and universities. Our products are also manufactured under the international ISO 9001:2000 Quality Management System.

Features & Benefits

- Data acquisition compliant with GLP and 21 CFR Part 11 regulations
- Centralized authorization system that supports an unlimited number of users
- A system that does not allow unauthorized access or tampering of files
- Public key Private key user validation to guarantee user authentication
- Easy installation, administration, maintenance and record keeping
- User-friendly system that does not hinder research



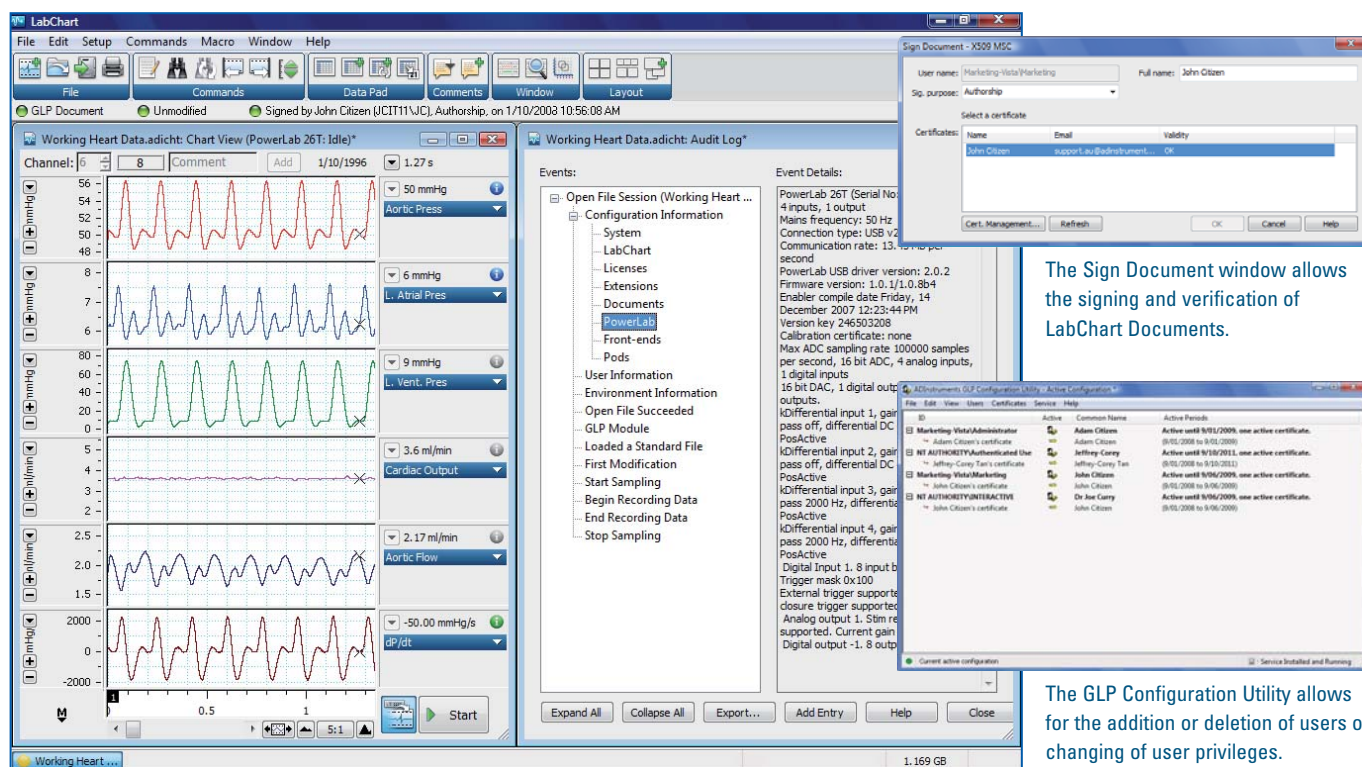
GLP Client and GLP Server Software

GLP Client Features:

- Secure signing of data files to eliminate unauthorized access and tampering (public key private key)
- Association of any changes to the data file with the registered individual making them
- A non-editable audit trail records all file operations
- Preservation of raw data
- Exportable (TXT, HTML, XML) and printable audit trail
- Incorporation of date and time stamping

GLP Server Features:

- Centralized user and signature authorization system
- Verification of signature validity when files are opened
- Use of digital signatures when saving data files
- Support for an unlimited number of users
- Access to remote configuration
- Extensive and customized logging
- Underlying configuration file suitable for scripting and automating batch processing



The Sign Document window allows the signing and verification of LabChart Documents.

The GLP Configuration Utility allows for the addition or deletion of users or changing of user privileges.

Above: The GLP Client adds a GLP Status Bar to the LabChart application window and an Audit Log (right panel) that records the user, GLP status, configuration and recorded data information.

For more information on how ADInstruments systems can expedite your research and facilitate GLP and 21 CFR Part 11 compliance, contact your nearest ADInstruments representative.

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PowerLab systems and signal conditioners meet the European EMC directive. ADInstruments signal conditioners for human use are approved to the IEC60601-1 patient safety standard and meet the CSA C22.2 No. 601.1-M90 and UL Std No. 2601-1 safety of medical electrical equipment standards.



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ISO 9001:2000 Certified Quality Management System

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