# InstantGMP™ PRO

## FOR REGULATED INDUSTRIES

InstantGMP PRO is the all-in-one manufacturing and quality system that delivers both efficiency and GMP documentation ... GMP-by-design...

INSTANTGMP™

# Our Electronic Batch Record centered system design stands out..... why?

- **It Provides Real-Time Resource Control...** facilities, equipment, inventory, and cost production info within the EBR framework
- **Make-To-Order Capability with New External User Batch Approval** provides product and process design-as-you-produce within the operation... ideal for both contract manufacturing and R and D development
- **Room and Equipment Log/Equipment Task Scheduler integration** for high-ticket resource control within the EBR
- **Two Level Configurability of EBR's...** the order and logic of the batch steps, and with our included Dynamic Fields toolkit, the batch steps themselves, for yields at a step, etc.
- **Project Level Operation in Real-Time** for white label, private label, etc. Dedicates EBR's, Resources and Production Teams
- **EBR's for Intermediates...** a completed batch can be an ingredient to another, for packaging operations or further work in process
- **Quality EBR Signoffs:** Quality has ability to issue, monitor and co-approve all EBR's to support GMP in the production process
- **EBR at the Center of the Design... by Design.** This vantage point allows better overall useability to optimize batch completion time, events, and costs

## Why Invest in InstantGMP PRO?

- Manage and Reduce Batch Time and Cost, while
- Generating GMP data Efficiently
- Work with a Team of Experts to fulfill a successful GMP Journey and Destination
- Minimize Batch Rejects and Reworks
- Secure, Manage, Stage, Track, Trace, and Ship Inventory End-to-End
- Track Costs Accurately with Configurable Costing and Inventory Cost Adjustment
- Reduce Errors with an Integrated System Footprint
- Streamline A Full Range of Quality Tasks with Included CFR Quality Workflows
- Use this Fully GAMP 5, Part 11, ISO 13485 Validated System with Confidence for all your regulated products
- Create Documents Interactively
- Version, Approve, and Deploy Documents and other files to Support a Full Range of PRO Operations
- Supported by an CBD/Hemp/Cannabis Experienced Regulatory team, top to bottom
- Schedule periodic equipment tasks and report completion status to select users

# GMP Inventory by design and detail

**Material Planning** makes it easy to procure ingredients by planned production times

**Cost the batch your way** Roll up materials with incoming costs... then configure labor, departmental time, etc... then price it!

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**Configurable screens and powerful inquiries** for securing and exporting real-time inventory data as you demand

**State of inventory color coding** identifies expired receipts or those needing retest

**Choice of inventory use...** barcode swipe, picklists for staging, selection form all approved receipts...all with max-min window for ensuring correct quantities

## Inventory Designed Bottom Up for GMP

- Specifications for materials vendor configurable for flexibility, linked to Material Receipt for convenience
- GMP role-based inventory status approval workflow built in
- Complete audit trail systemwide
- Validation per GAMP and FDA standards for any type of GMP regulated product
- Configurable FG costing provides accuracy and even creativity for a variety of cost types
- Shipping Manifest closes the GMP loop to the customer or end user

## System Generated Inventory Reports for Better Inventory Decisions

- Current Inventory
- Inventory Value
- Low Inventory
- Vendor Lot Traceability
- Inventory in Quarantine
- Production Lot Traceability
- Alert Level
- Reorder Level
- Materials Near Retest/Expiry

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# Document Management

The InstantGMP™ PRO Electronic Document Management System provides easy document approval, tracking, and versioning to support the GMP activities throughout the PRO system.

- **Document Control:** Document managers can upload, version up, and route documentation for review and approval.
- **Document History:** The system maintains all superseded documentation alongside the comments and changes made during an activity cycle.
- **Documentation for Active Support:** The EDMS is integrated into most of the software. This means key documentation can be attached directly to a Batch Record, a Room or Equipment Log, or a material specification.
- **Wide variety of file types supported**

## Documentation Included with Purchase:

**120 Template GMP Policies and Procedures for areas like:**

- Quality Systems
- Materials
- Facility & Equipment
- Safety
- Packaging and Labeling
- Production

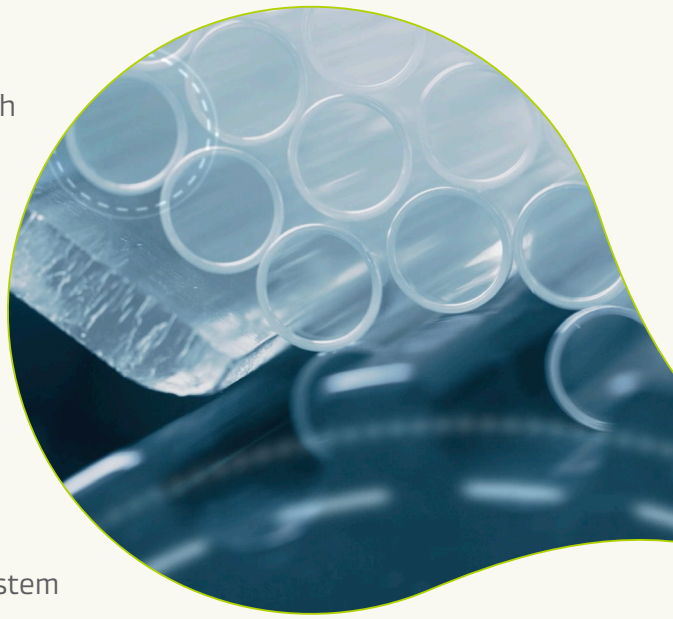
**18 GMP Forms and Templates:**

- Audit Plan
- 2-Way CDA
- Deviation Report
- Serious Adverse Event Report
- and much more!

# Quality Management

InstantGMP™ PRO can also help your quality team manage a complete set of tasks in a cGMP operation.

- **Quality Logs:** Manage the many different quality logs such as: **Deviation**, **Incident**, **CAPA**, **Change Control**, and **Audit**. With pre-built workflows to make navigation and completion flow easy
- **Integration of EBR and Deviation Workflow** to save time in documentation and resolution
- **Vendor Management:** Collaborate on key documentation with your vendors in "data room" mode to complete and expedite vendor qualifications, agreements, etc.
- **Training Log:** Issues and stores training activities completed by your personnel. Used in conjunction with the optional **Learning Management System**, supplied by Entrenar.se, organizes online curricula that can include presentations, videos, or documents for all included users on the system.
- **Vault:** The Vault gives the end-user an internal file database and organizer separate from the Document Management System to secure the hierarchy of non-versioned information.



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