

InstantGMP™ Recognized Among Top 8 GMP Software Solutions for 2025 by SafetyCulture

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Dr. Richard Soltero

InstantGMP™, a leading provider of manufacturing and quality management software for regulated industries, proudly announces its selection by SafetyCulture as one of the Top 8 GMP Software Solutions. The honor places InstantGMP alongside other elite platforms dedicated to streamlining Good Manufacturing Practices (GMP) through innovative digital tools.

As featured in SafetyCulture's Top 8 GMP Software of 2025 list, InstantGMP continues to set the standard in the field

by offering a powerful, all-in-one solution that supports GMP-certified operations across pharmaceutical manufacturing, dietary supplements, biotech, and 503B compounding pharmacies.

"It's a privilege to be recognized by SafetyCulture as one of the best GMP software providers for 2025," said Dr. Richard Soltero, President of InstantGMP. "We remain committed to providing manufacturers with intuitive tools that simplify compliance, reduce risk, and improve quality throughout the entire production lifecycle."

InstantGMP™ PRO was highlighted in the report for its modular, cloud-based design that helps facilities manage every aspect of GMP, from training to production documentation, with unparalleled efficiency.

The platform integrates:

- ☐ Electronic Batch Record Software (EBR): Streamlines the creation, execution, and review of batch manufacturing records (BMR), including master batch records (MBR), ensuring accuracy, traceability, and data integrity.
- Inventory Management: Innovative manufacturing inventory software that enables real-time

material distance, from fatt more exercise and again man produce to lease.
 Quality Management: User-friendly tools to monitor deviations, manage CAPAs, and verify
GMP-certified practices across the facility.
 Learning Management: Assigns and tracks required GMP training, reinforcing regulatory
readiness and employee competence.
☐ Document Management: Securely stores SOPs, specifications, and audit records for easy
access and compliance during inspections.
☐ Equipment Log: Tracks equipment use and maintenance in real time to support preventive
care, reduce downtime, and improve overall equipment effectiveness, all within a GMP software
environment.

material tracking from raw ingredient receipt through final product release

Each module is pre-validated to save time and ensure compliance, making InstantGMP an ideal fit for small to mid-sized enterprises looking to adopt manufacturing process software without the complexity of multiple systems.

InstantGMP's platform is uniquely equipped to support regulated facilities in maintaining and documenting compliance with FDA regulations and evolving GMP standards. In addition to comprehensive batch management tools, the software simplifies the creation and execution of detailed records such as master batch records (MBRs) and batch manufacturing records, giving teams complete control over production documentation and traceability.

Whether managing a 503B compounding pharmacy or scaling biotech operations, InstantGMP delivers a seamless experience with out-of-the-box features, intuitive interfaces, and exceptional customer support for manufacturers in English and Spanish-speaking countries.

Contact us today to discover how our GMP software can help you create and manage batch manufacturing records, streamline inventory with powerful manufacturing process software, and stay GMP certified with confidence.

About InstantGMP™, Inc.

Founded by pharmaceutical industry veteran Dr. Richard Soltero, InstantGMP, Inc., offers affordable all-in-one manufacturing, inventory, and quality software. The company develops cloud-based electronic batch record software and standard operating procedures specific to industries that are required to follow FDA manufacturing regulations and Good Manufacturing Practices ("GMP").

As a manufacturing software company, InstantGMP™ pioneered accessible, easy-to-use electronic batch record software for products manufactured using GMPs. The Company's updated software simplifies the documentation and approval procedures for quality processes that keep all quality documentation organized in electronic format while providing for quality checks and workflow processes to make compliance with FDA requirements easy.

Debbie Young
InstantGMP
+1 919-645-1073
email us here
Visit us on social media:
LinkedIn
Facebook
YouTube

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