InstantGMP Expands 5-Point Implementation Plan with On-Site Services That Deliver Faster ROI for Manufacturers

Expanded 5-Point Implementation Plan

CARY, NC, UNITED STATES, November 25, 2025 /EINPresswire.com/ -- InstantGMP™, a leading provider of GMP-compliant manufacturing software for pharmaceutical and regulated industries, announces an enhancement to its 5-Point Implementation Plan with expanded on-site implementation services designed to help customers achieve measurable return on investment (ROI) faster, streamline system adoption, and accelerate GMP and FDA compliance readiness.

The recently launched 5-Point Implementation Plan provides a structured, strategic roadmap for onboarding customers to the InstantGMP™ PRO manufacturing execution and electronic batch record system. Built on years of customer feedback and industry experience, this plan ensures every organization receives guided support, expert training, and the tools needed for long-term operational success.

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>
> *Brandy Irons*

Now, InstantGMP is elevating this framework by highlighting the powerful advantages of on-site implementation, particularly for manufacturers seeking faster project momentum, reduced rework, and improved accuracy during setup.

InstantGMP’s on-site approach helps teams move from planning to execution with greater confidence, speed, and clarity.

Key benefits include:

Increased Project Commitment and Faster Launch Timelines

When customers schedule an in-person implementation visit, their internal teams are more likely to dedicate focused time to onboarding tasks. This prevents delays and helps ensure the project stays on track instead of being repeatedly postponed for competing operational priorities.

Tailored Training for Different Learning Styles

While virtual calls are effective, some teams learn best through hands-on instruction. Being on-site allows InstantGMP trainers to teach and immediately execute software processes in real time, resulting in stronger comprehension and faster mastery.

First-Hand Understanding of Facility Workflows

Seeing a facility's operations, production flow, and equipment layout enables InstantGMP to better understand customer challenges. This ensures system configurations are correct the first time, reducing the need for rework, re-entries, or post-launch adjustments.

Real-Time Collaboration With Undivided Attention

On-site visits ensure that the InstantGMP and customer teams can both ask questions, validate steps, and resolve issues immediately. This two-way communication accelerates progress and eliminates bottlenecks that often occur during virtual-only collaboration.

Faster Setup of Complex Manufacturing Records

Certain implementation tasks, such as building initial Master Production Records (MPRs) and Master Test Protocols (MTPs), can be completed significantly faster by InstantGMP experts when they are physically on-site. This is especially impactful for formulas requiring intricate calculations or multi-step routing, allowing these tasks to be completed in parallel while customer staff focus on other project components.

"Being on-site gives us a clearer understanding of each customer's unique processes, which means we can configure their system correctly the first time," said Brandy Irons, Senior Training and Technical Support Manager/Quality Assurance at InstantGMP. "It also strengthens training, communication, and collaboration, which in turn helps teams get up and running faster with InstantGMP."

These enhanced on-site services seamlessly integrate with InstantGMP's existing 5-Point Implementation Plan, which includes:

Phase 1: Planning & System Design
Phase 2: System Configuration & Implementation
Phase 3: Pilot Testing, Operator Training & Launch
Phase 4: Optimization
Phase 5: Ongoing Configuration Assistance

Together, these phases provide a complete, expertly guided implementation strategy that transforms system adoption into a structured, measurable process aligned with GMP and FDA compliance requirements.

To learn more about InstantGMP's enhanced on-site implementation services and 5-Point Implementation Plan, contact our sales team today.

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.

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