

It takes weeks or months to release a vaccine batch to the FDA for review. This big pharma customer previously had a complex batch record consisting of over 1,200 pages and taking over 30 days to complete. Every page had multiple opportunities for human error including incorrect entries, miscalculations, and executing steps out of order. Administrative controls as well as rigorous and time-consuming batch record reviews were required at the end of production to ensure product quality.



By implementing an electronic batch record in MES, this pharma manufacturer practically eliminated opportunities for error—enabling more predictable batch release cycle times, and ultimately reducing finished goods inventory.

Continua enables process manufacturers to move up the Digital Plant Maturity Model through the application of innovative process automation, workflow, and data management solutions. Its team of expert consultants help companies deliver tangible business value through increased efficiencies and by leveraging new and emerging data architectures and associated technologies.

The business primarily serves regulated manufacturers who face growing pressures to deliver more product at a lower cost, while ensuring quality and security of supply.

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