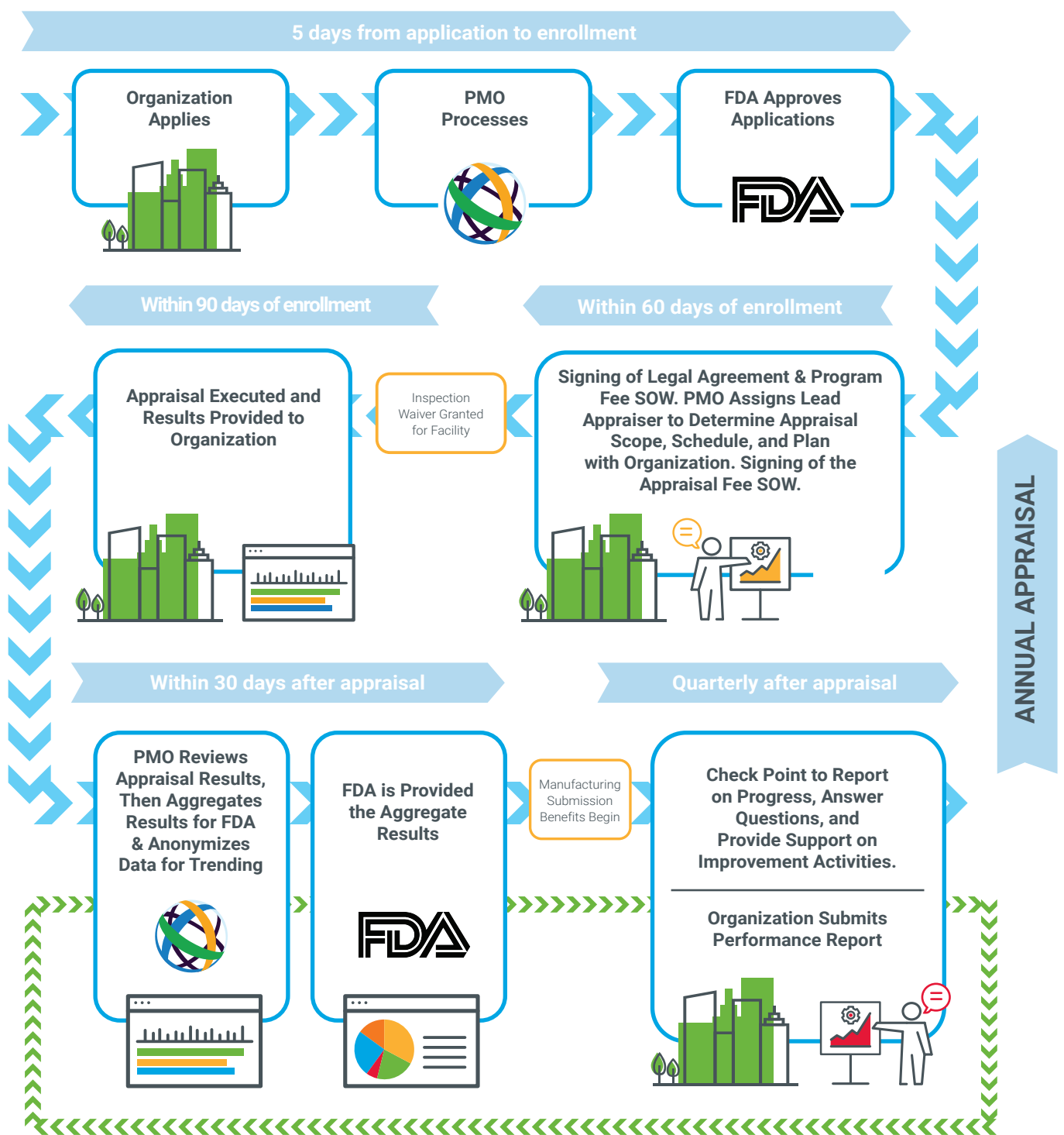


Innovating Better Medical Devices Faster at Lower Costs

Medical device manufacturers with a strong compliance profile can sign up to be industry leaders in a pilot program that aims to offer better patient safety outcomes through performance improvements against a set of proven best practices. Results may pay for themselves through a competitive advantage of increased quality, lower costs, and saved time.

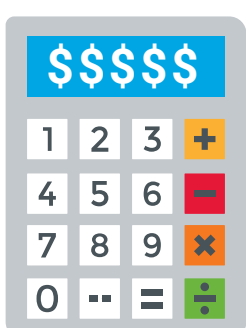
How Does MDDAP Work?

This is not an invitation to heightened regulatory scrutiny by the FDA—there is no downside to participating in the pilot. **This voluntary “conversational” engagement is focused on understanding how to improve quality and outcomes;** it is not focused on “checking the box” for regulatory compliance.



Program Cost vs. Return on Investment (ROI)

A standard \$5,000 annual Program Fee is required to support each enrolled facility in the program. A separate Appraisal Fee will be determined based on the scope of work required to complete the appraisal and varies based on company size, location, and complexity. Note, reduced fees are available for organizations registered as a “small business” with an FDA SBD number. All enrolled organizations stand to benefit from a cost, time, and resource perspective.



One organization realized annual savings of nearly \$300k

Another organization increased production capacity by 11%, resulting in an additional \$15M in sales