



The endpoint™ difference

Global presence

With hundreds of active trials under way around the clock and in locations around the globe, endpoint is helping biotechnology companies and contract research organizations experience the power and simplicity of PULSE™, our proprietary technology platform.

Deep experience

endpoint was founded in 2009 and our leaders have been developing interactive response technology (IRT) systems since 1998. The endpoint team brings expertise in IRT technology, clinical trial study design and supply chain management to the design and development of IRT solutions.

Rapid, customized IRT

PULSE enables design, modification, testing and launch of a fully customized and validated IRT system, typically in 30 days, and at less cost than traditional IRTs. Enhancements can be made easily without impacting your study.

Flexible technology

Our highly configurable and modular technology platform relies on reusable components that provide greater flexibility than legacy IRT systems and without any sacrifice in quality.

Comprehensive solutions

PULSE delivers broad functionality that meets the demands of clinical study managers. We can design and deploy a tailored IRT system that gives you the precise features you desire and access to data and study management capabilities from multiple platforms, anytime and anywhere.



Meet PULSE™ The new standard in IRT systems.

PULSE is a highly configurable platform with an intuitive user interface that enables the rapid development of customized IRT systems for clinical trials. As a state-of-the-art technology, PULSE

delivers unprecedented speed, flexibility and control.

Mobile

Stay connected to your clinical trials via smartphone or tablet. A mobile-optimized interface gives you access to IRT system functionality, meaning you won't miss a

beat, no matter where you are.

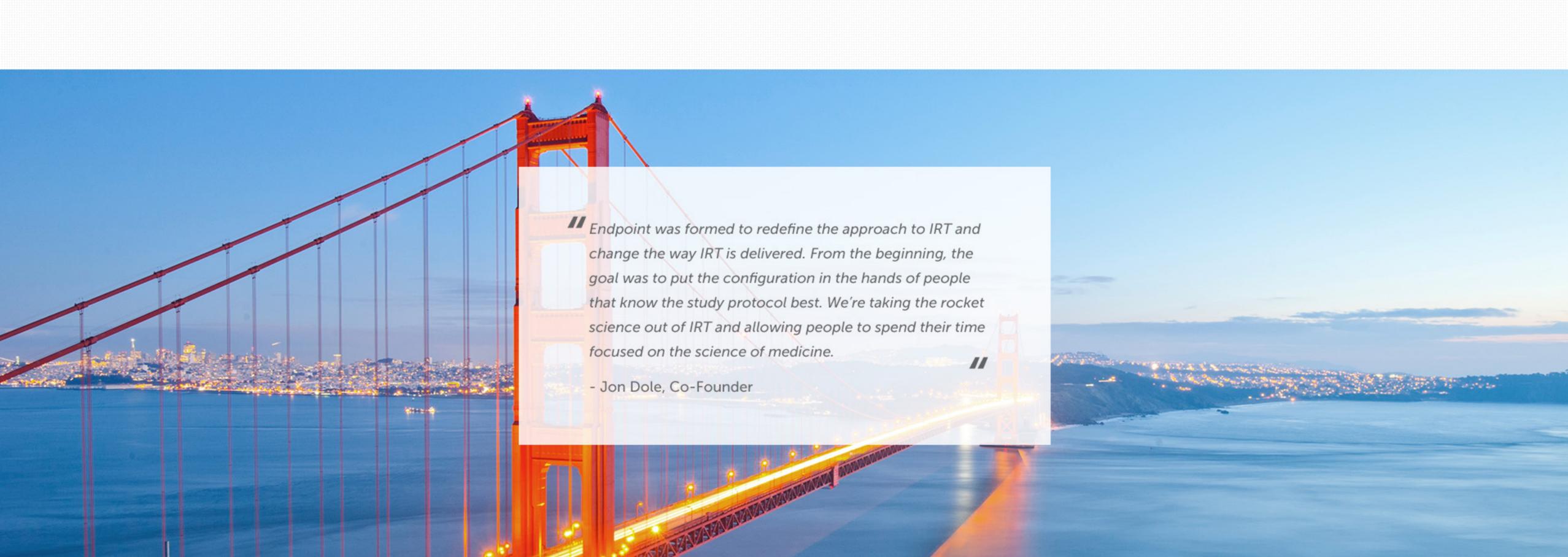
SaaS

Our software-as-a service (SaaS) model allows you to design and deploy your IRT systems in the cloud as hosted solutions. 2

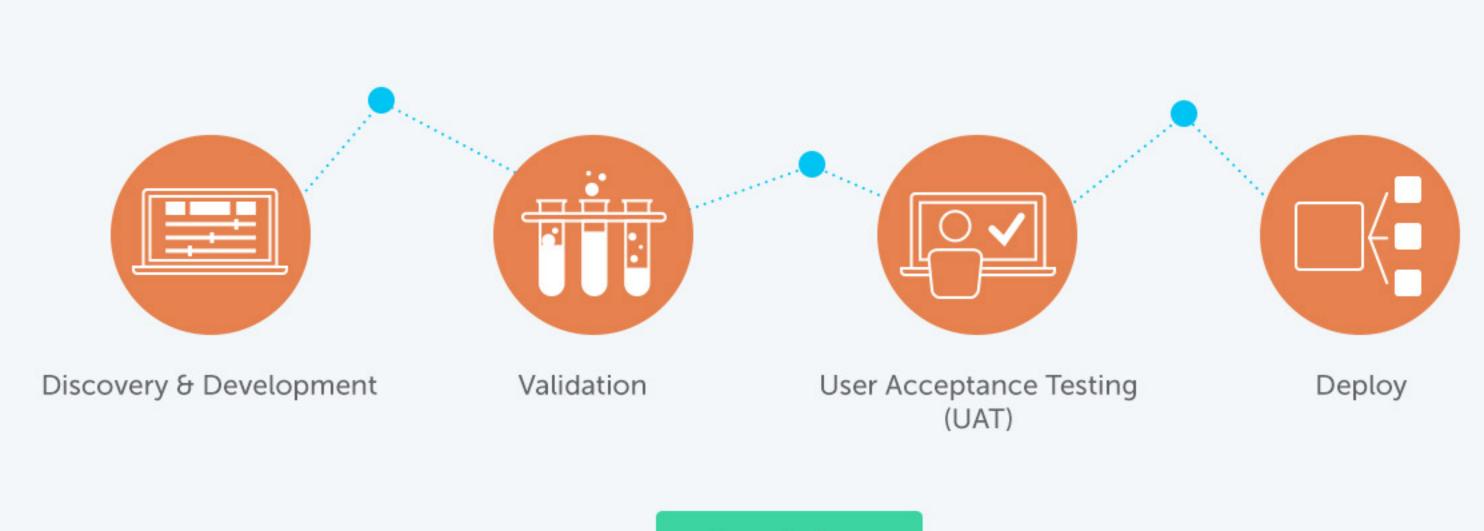
Integration

PULSE integrates seamlessly with other systems, including clinical supplies, EDC, ePRO and CTMS.

Learn more

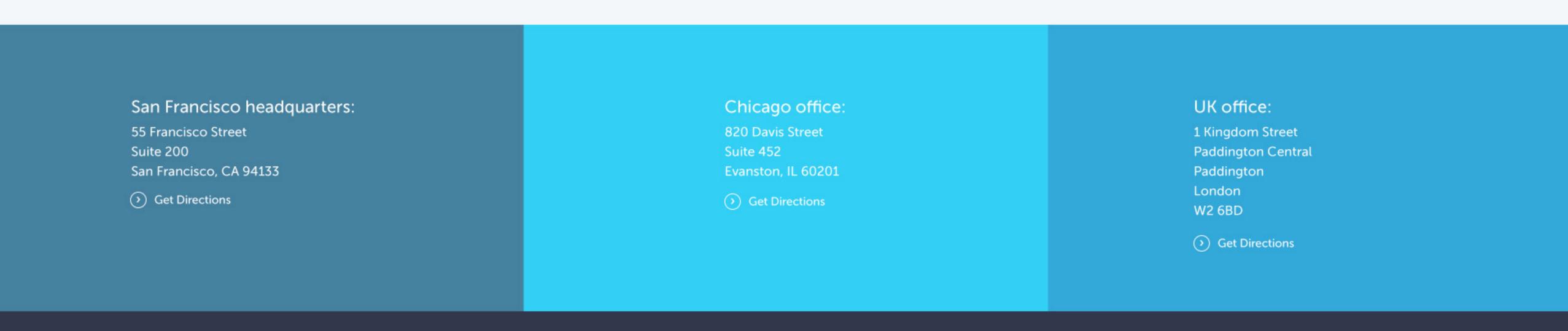


30 days to 'study start' date



Request a demo

Office Locations



Connect with us **in**

endpoint clinical

Faster to deploy, easier to manage, higher quality than ever before.

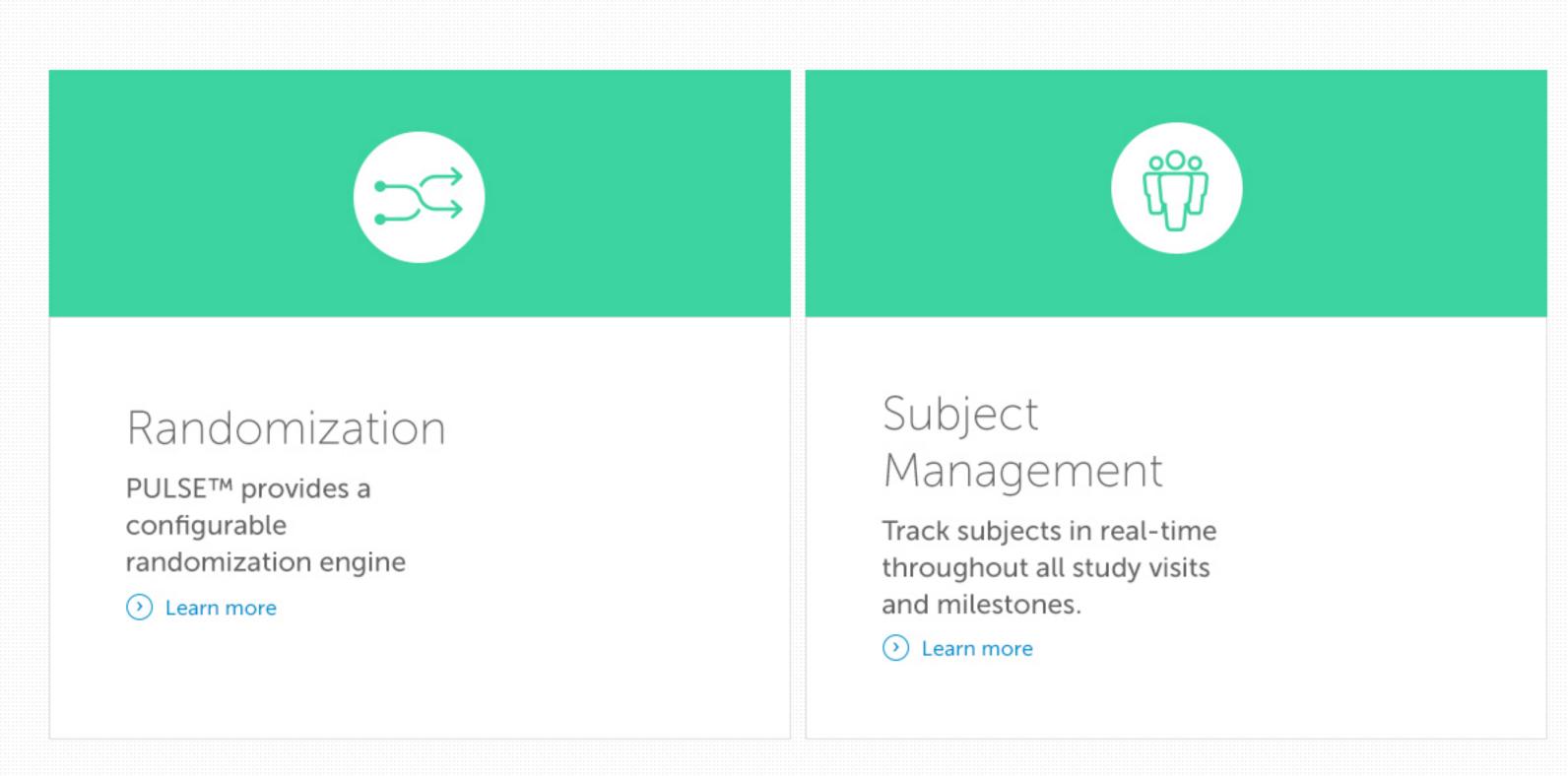


It's not rocket science...anymore.

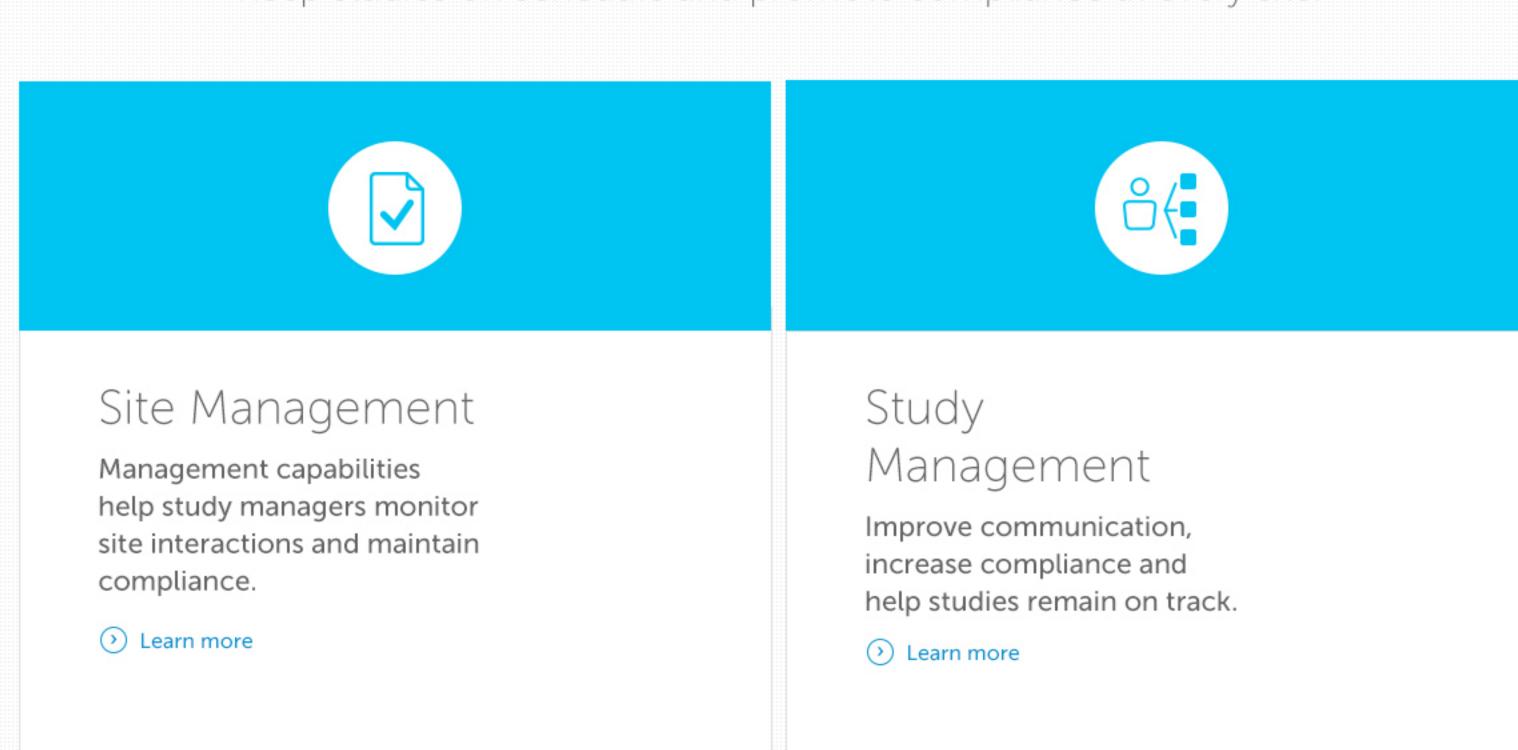
endpoint has completely redefined the integrated response technology (IRT) system by developing a unified platform. With endpoint PULSE™, our proprietary technology, pharmaceutical and biotechnology companies and contract research organizations (CROs) can access clinical study data from any platform—phone, web or mobile device—and segment data according to their specific needs. endpoint supports all aspects of clinical studies, including randomization and subject management, site and study management, drug supply management and ePro.

We partner with clients to design, develop, validate, test and deploy IRT systems. endpoint also licenses PULSE as a Software as a Service (SaaS) solution to certified users, enabling clients to configure and deploy their own systems.

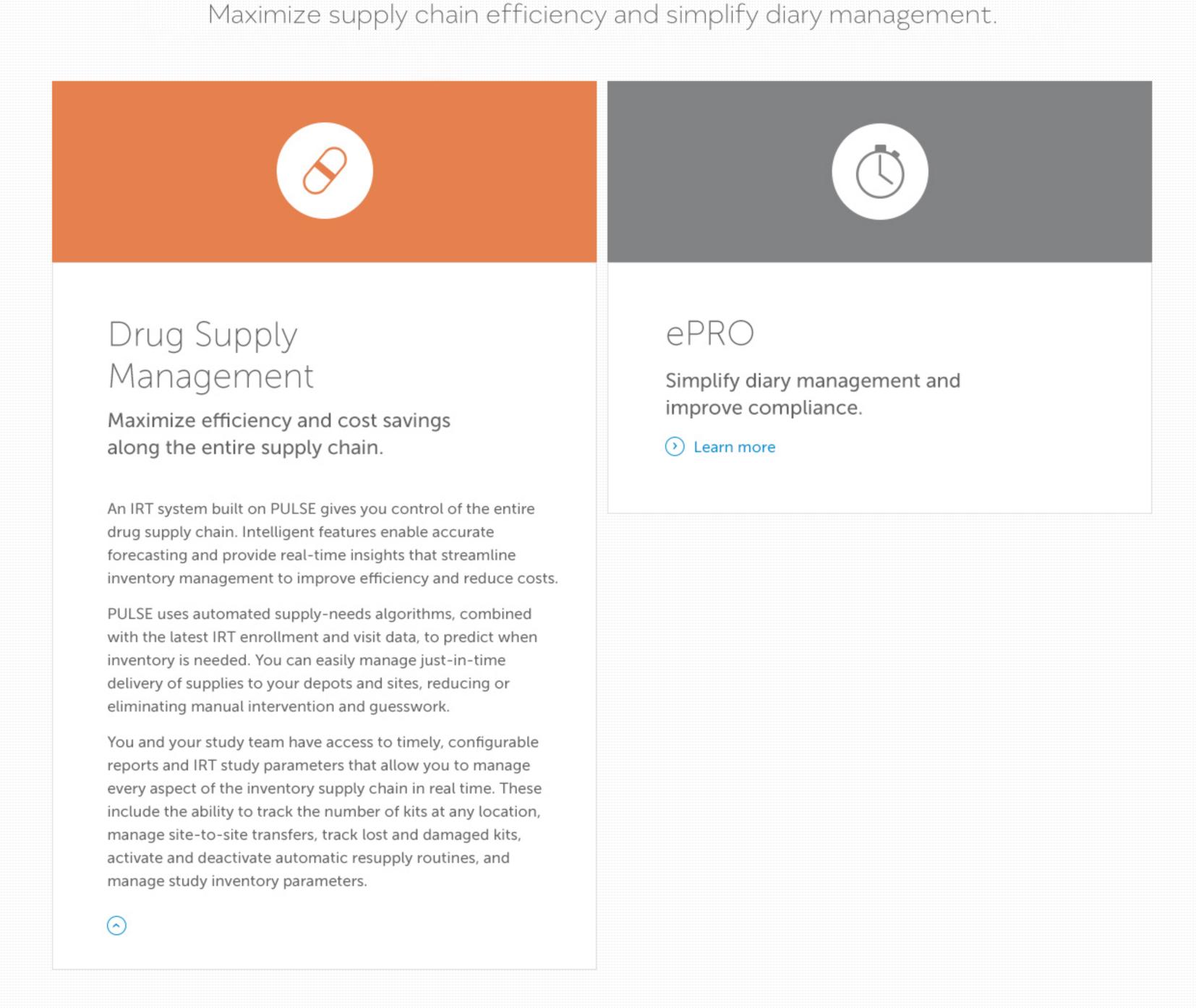
Randomization and Subject Management A next-generation platform to support any trial.



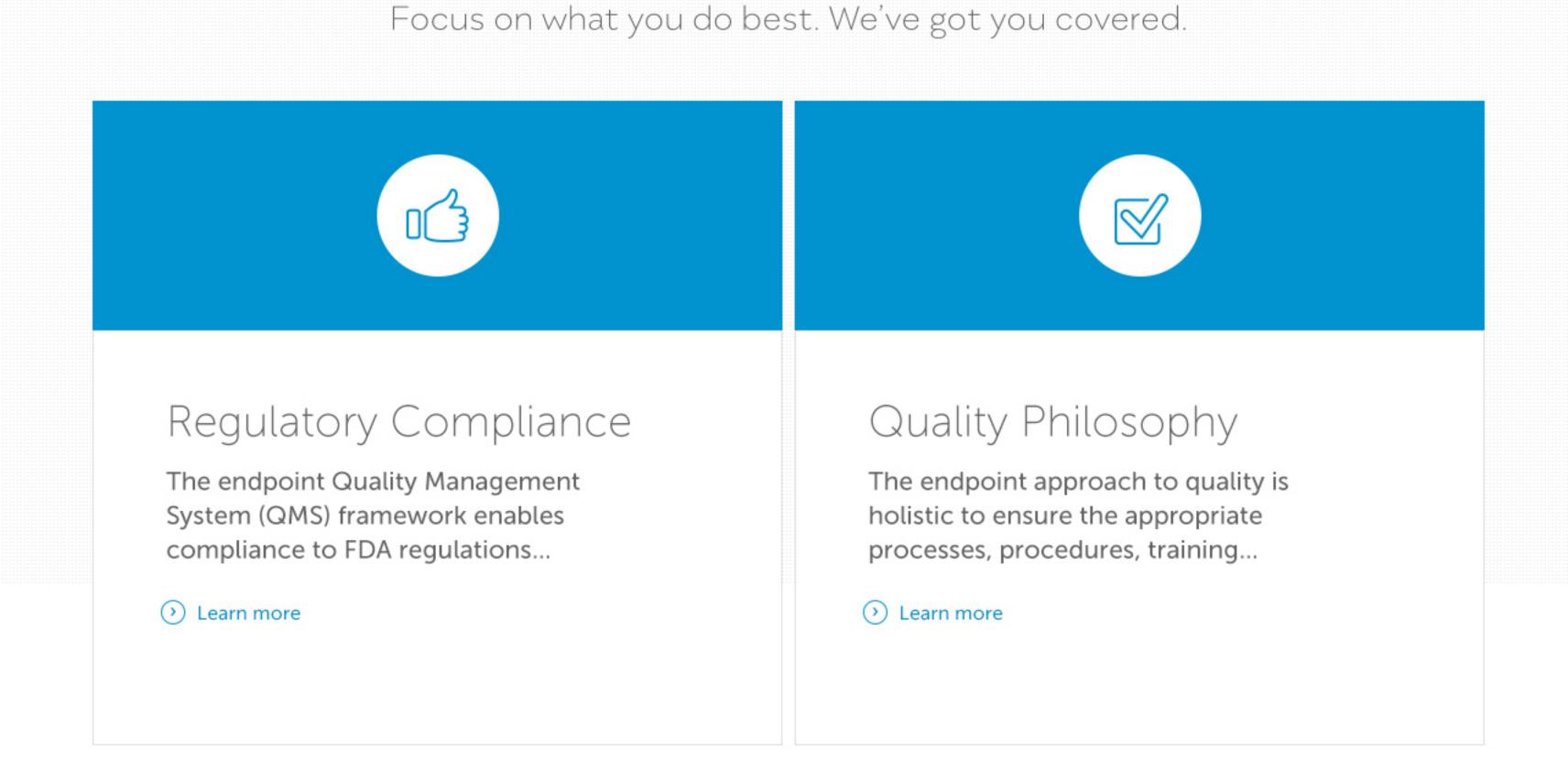
Site and Study Management
Keep studies on schedule and promote compliance at every site.



Drug Supply Management and ePro



Compliance and Quality



Ready to get started?

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Office Locations

San Francisco headquarters:
55 Francisco Street
Suite 200
San Francisco, CA 94133

O Get Directions

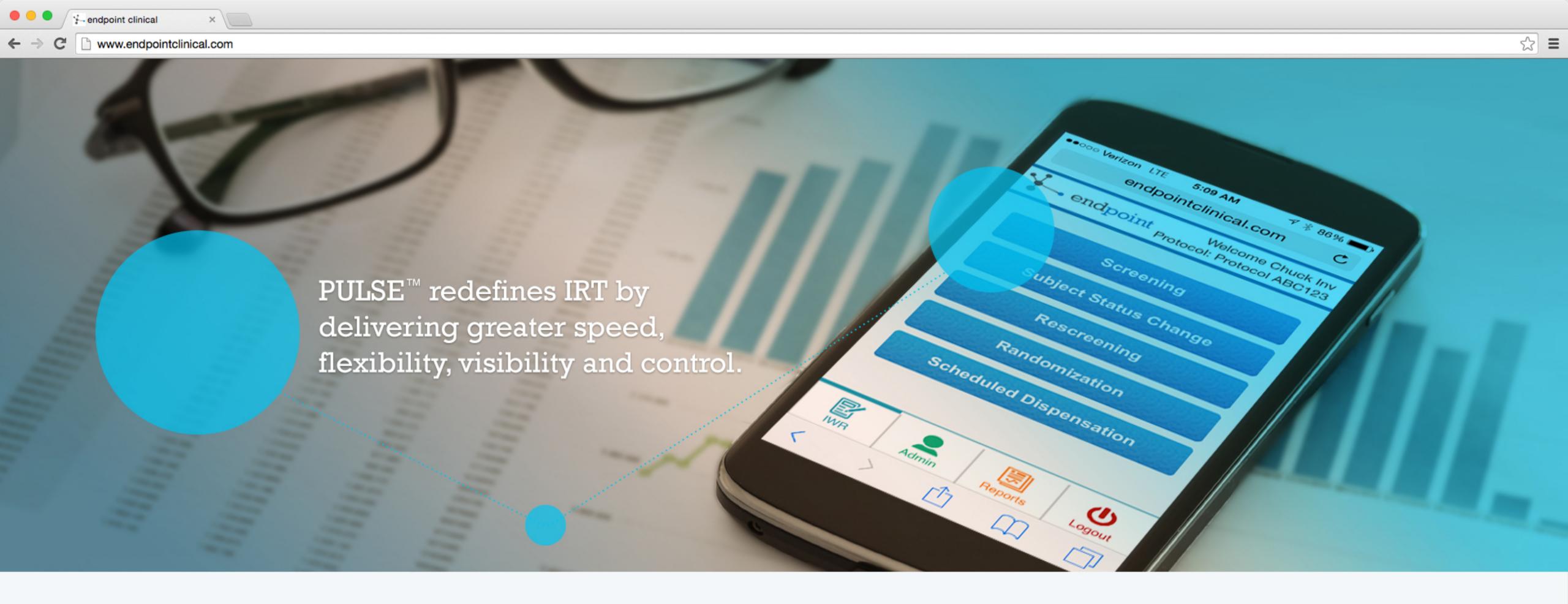
Chicago office:
820 Davis Street
Suite 452
Evanston, IL 60201

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UK office:

1 Kingdom Street
Paddington Central
Paddington
London
W2 6BD

3 Get Directions

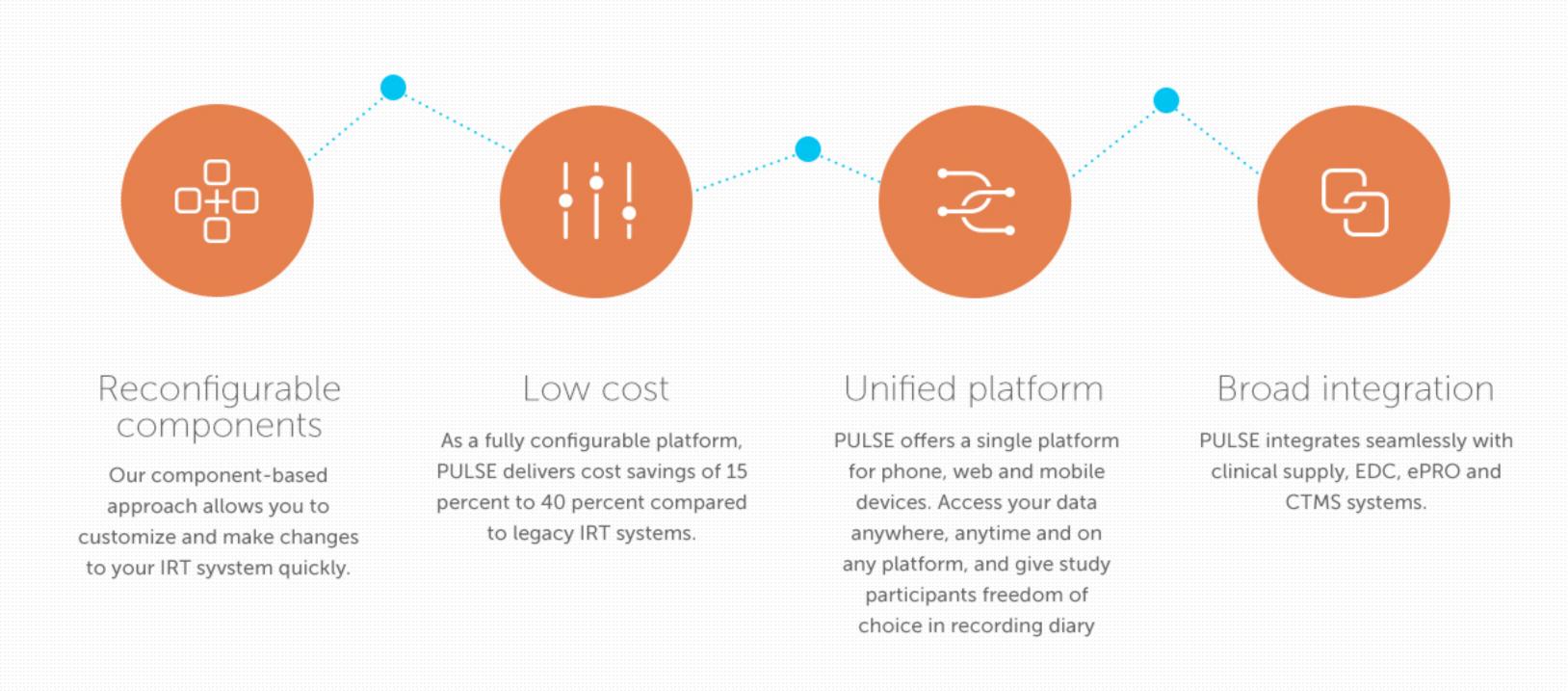


Taking IRT systems to the next level.

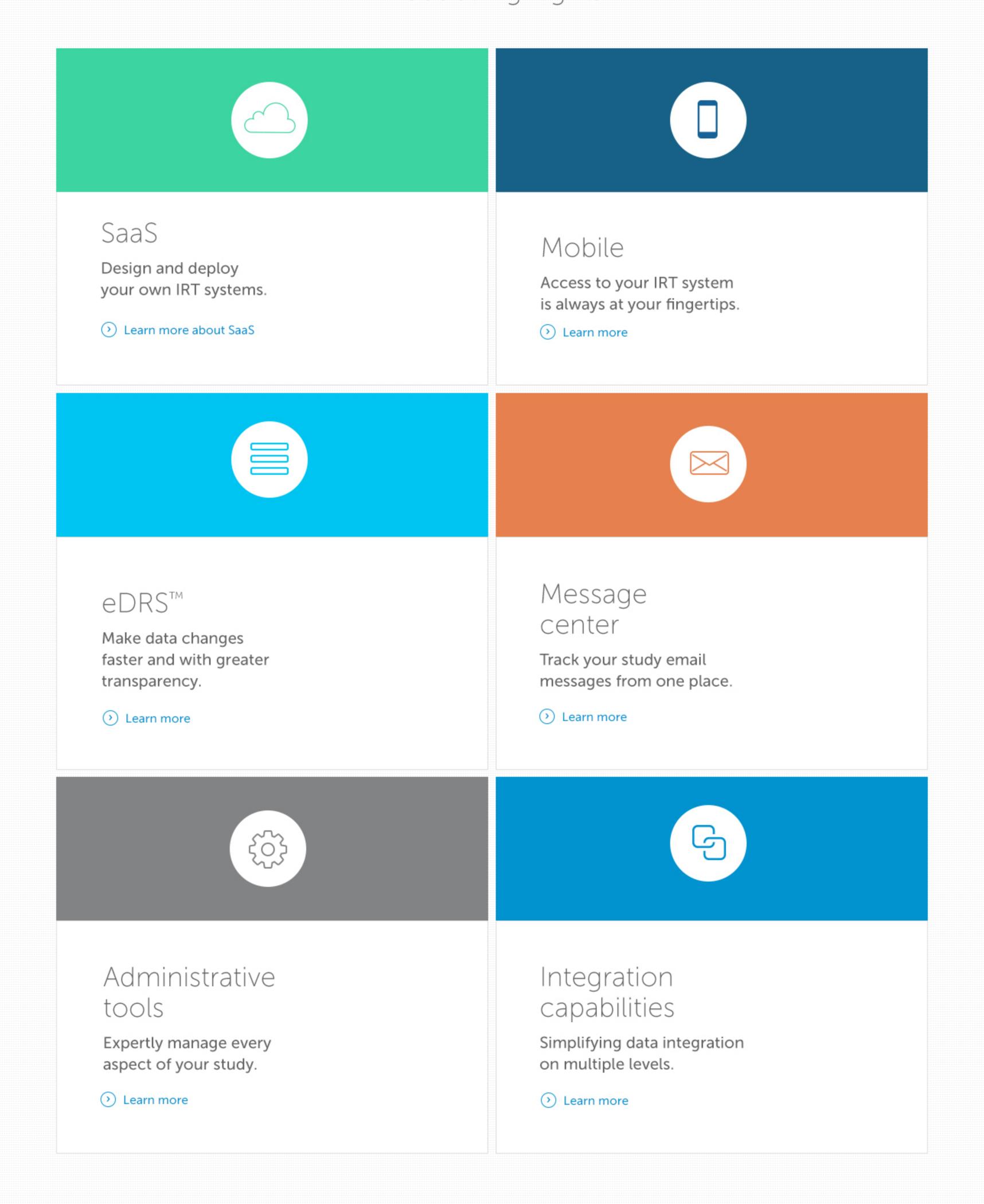


The PULSE platform enables rapid development of a fully customized and validated IRT system to meet all your trial design needs—at a much lower cost than with other providers and in a fraction of the time. We can help you develop a customized system in 30 days, much faster than the industry standard of eight to 16 weeks, and without any sacrifice in quality. An IRT system built on PULSE streamlines clinical trial randomization; drug supply management; and site, study and subject management.

PULSE is endpoint's proprietary rules-based core engine. Pre-validated, configurable study components provide flexibility to enable rapid development and modifications to IRT systems.



Product highlights



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