STUDY PLANNING AND SITE OPTIMIZATION

Investigator/Site Training Platform

Exceptional Site Performance & Satisfaction Starts with an Exceptional Training Experience

ACCELERATED TIMELINES	REDUCE COSTS	IMPROVE QUALITY	INSPECTION READINESS	SPONSOR OF CHOICE
Gain 55 enrollment days through just in time training	**Cut training costs by over 60%** compared to traditional delivery	**Reduce** number of **protocol deviations**	**Avoid the #1 finding** of FDA Audits	Reducing the **burden of Site training by 50%**

Highly Effective

Sites have a clear understanding of requirements and an improved ability to adhere to protocol and trial amendments

Automated Processes

A highly automated system that deploys training quickly, with complete, real time tracking.

Reduced Burden

Reduce the burden on Site staff of unnecessary and/or redundant training

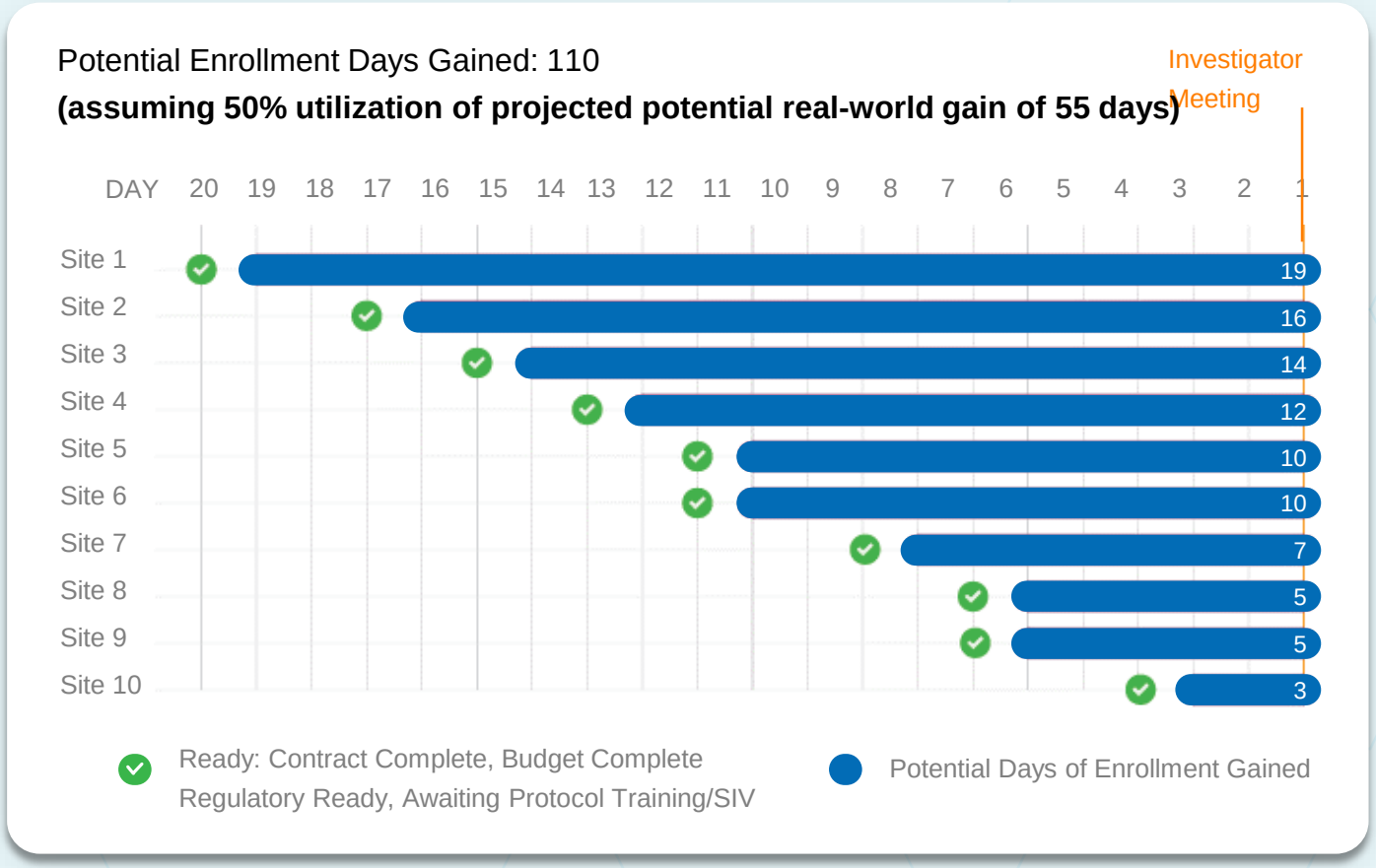
Gain time for enrollment through just in time training

ACCELERATED TIMELINES

With an on-demand training rollout, protocol training is available as soon as the site is ready (contract complete, budget finalized, and regulatory ready).

Rapid Training Development & Deployment

Implement your training program **in 5 working days.**



Sponsors save over 60% or More in Costs with Training Available to Staff 24/7

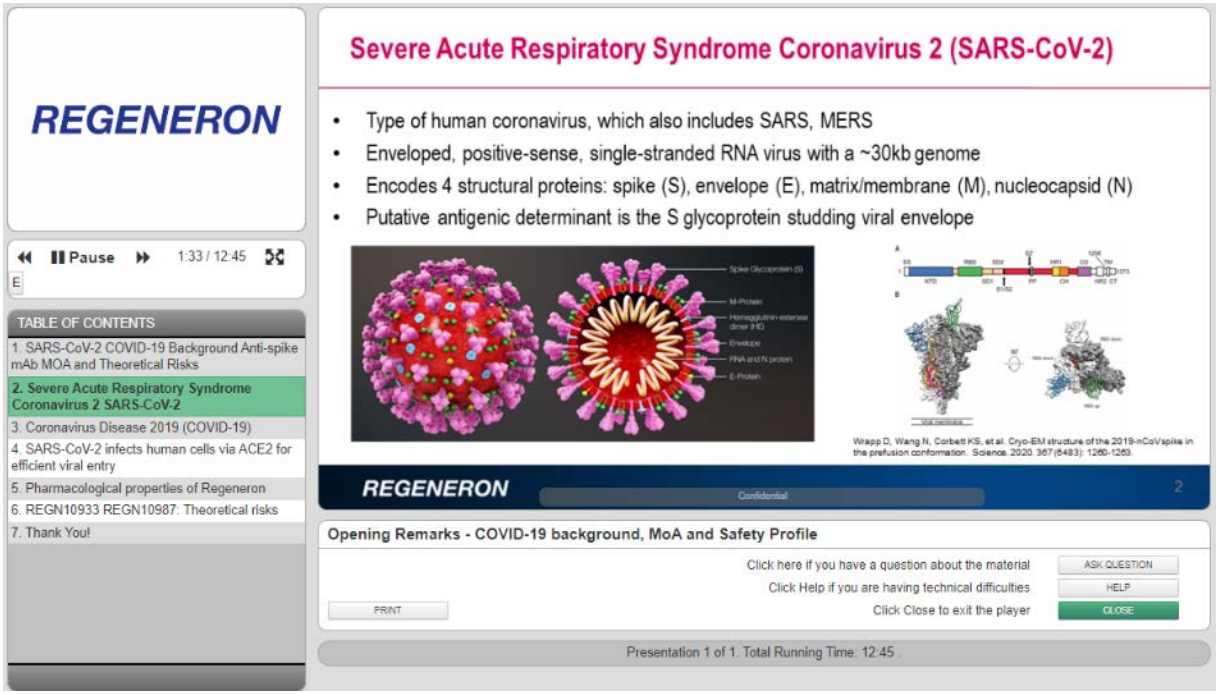
REDUCE COSTS

The ROI for using Trifecta is easily realized based on face-to face investigator meeting costs, but the financial benefits go beyond that:

- Estimated Cost to Attend Investigator Meeting in 2020 - **$2,400 per PERSON**
- Average Investigator Training Cost per site for Trifecta - **$2,500 per SITE**

Reduction in Sponsor/CRO costs, for example:

- **Reduction in overall Monitoring Costs due to less time on site for CRAs**
- **Reduction in Study Management time spent in audit preparation**

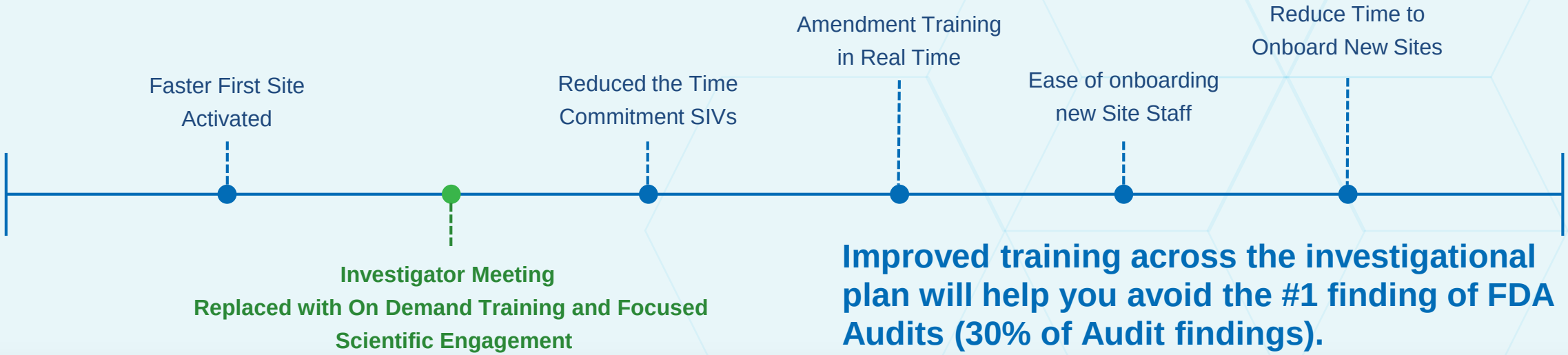


Training is available to site staff throughout the study

Improve Quality With Training That's More Accurate, Immediate, & Easy to Maintain

IMPROVE QUALITY

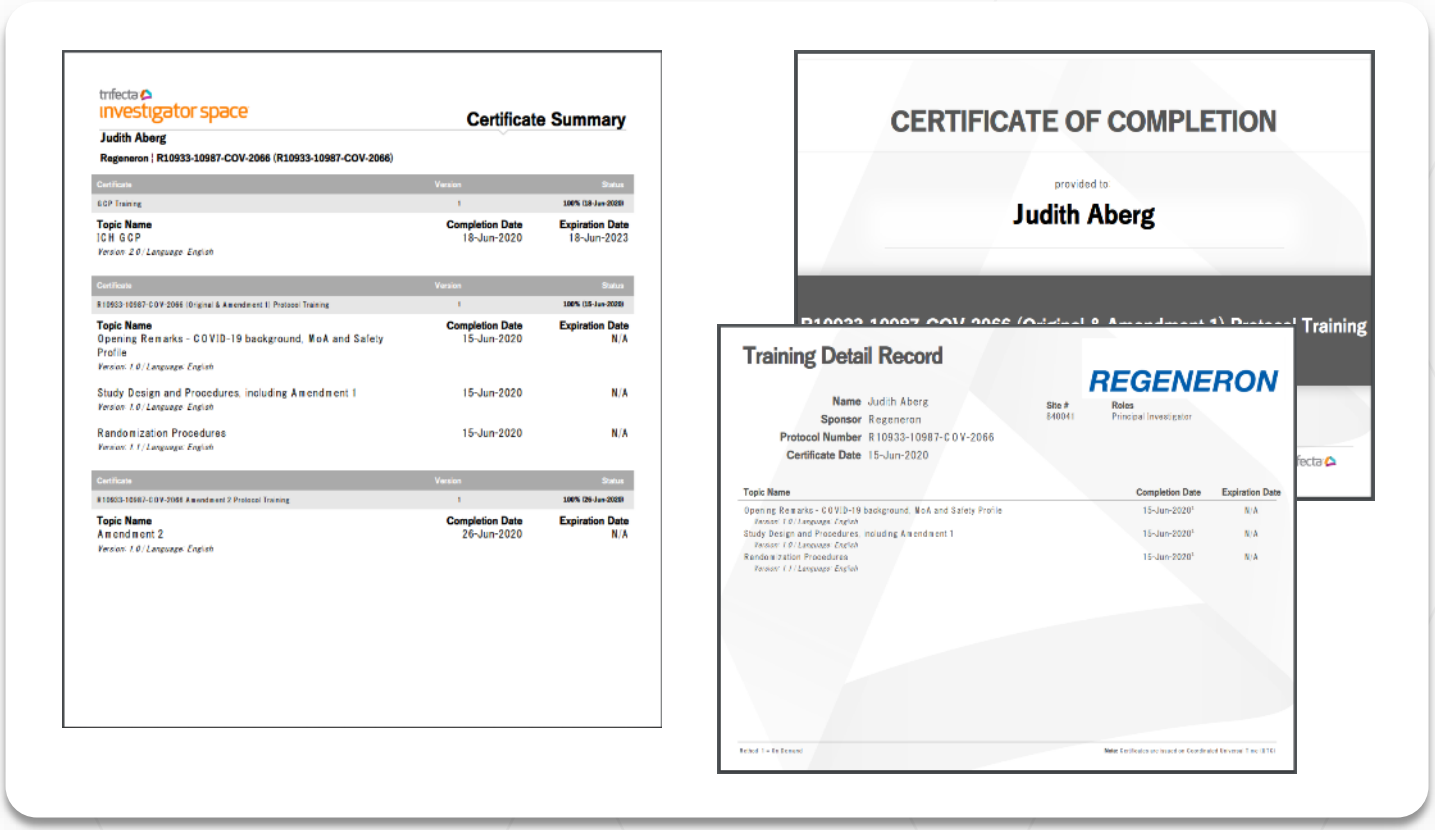
- Deliver "Just-in-Time" training across the study lifecycle
- Proprietary video-based training production process reduces asset creation time by 90%
- Updates & Amendments produced and available in 24 hours
- Immediate transformation of lessons learned into actionable guidance reduces data queries and data management/monitoring time
- Easy remediation of protocol deviations



Improved training across the investigational plan will help you avoid the #1 finding of FDA Audits (30% of Audit findings).

Be In a Perpetual State of Inspection Readiness

- ✓ Clinical Qualification Management System automates distribution, collection, and maintenance of all training documentation
- ✓ 100% security, auditability, & traceability
- ✓ Real-time compliance reporting provides total training process transparency

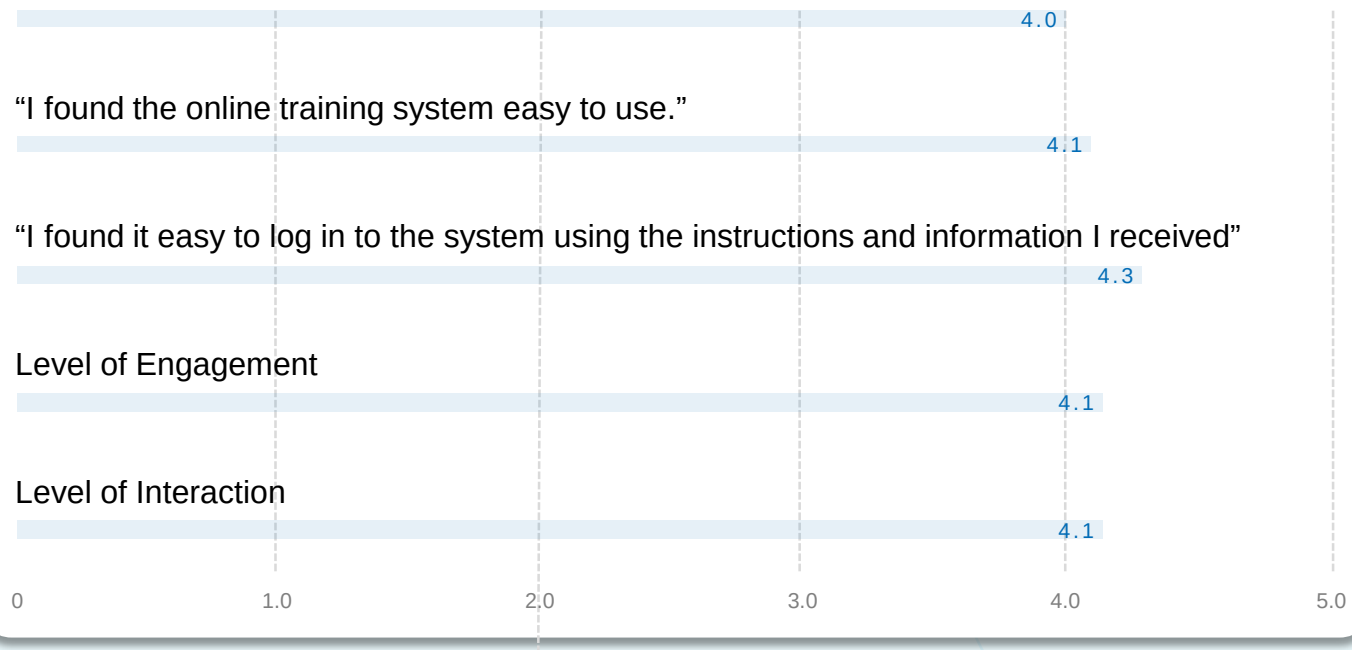


Improve Site Satisfaction, Eliminate Redundant Training

SPONSOR OF CHOICE

N=3,866 responses

Exceptional Feedback on Trifecta Training in 2020



STRONGLY DISAGREE — **STRONGLY AGREE**

Site Investigator Training Requirements

STUDY 1
Protocol W32
Mechanism of Action
SCORAD
GCP

STUDY 2
Protocol M91
GCP
SCORAD
eClinRO

STUDY 3
Protocol ZYY
Imaging Criteria
COX-2 Inhibition
GCP

STUDY 4
Protocol 45W
GCP
SCORAD
eCOA Tablet

Training Taken in Study 1 Automatically Credited for Any Additional Studies

In one year, more than **50% of Investigators** were exempt from redundant GCP training.