

Veeva Clinical Platformが実現する エンドツーエンドの試験管理ソリューション

11Nov2025



Vision

Building the Industry Cloud for Life Sciences

Values

Do the Right Thing

Customer Success

Employee Success

Speed



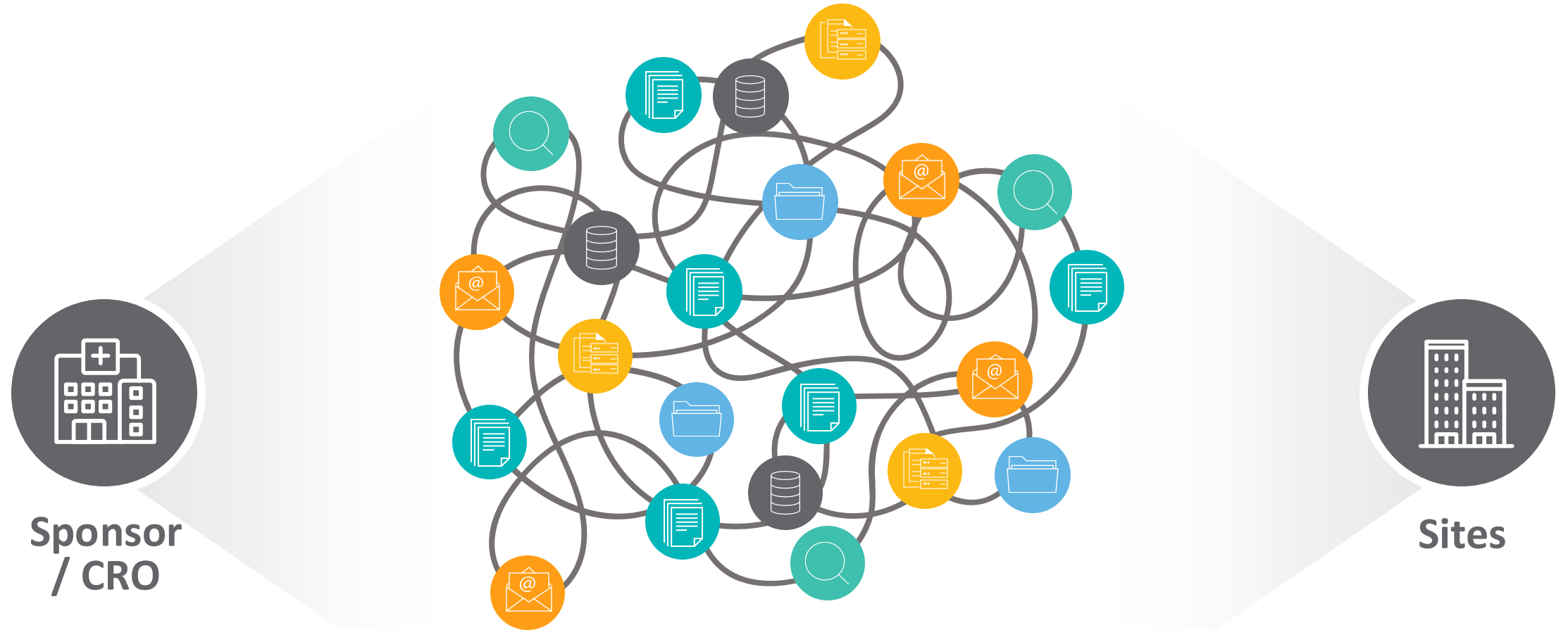
A Public Benefit Corporation

Veeva Clinical Vision



Simplify and Standardize
clinical trials for higher efficiency and a better
experience for sponsors, sites, and patients

Complicated Site Engagement Ecosystem



Multiple Systems

Information Silos

Manual Processes

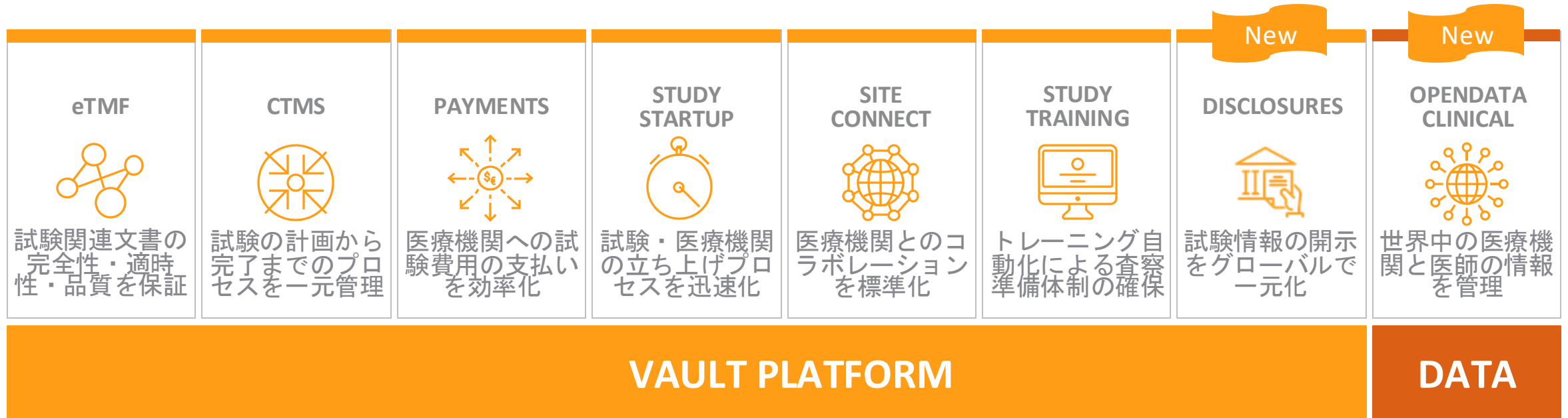
Content Replication

Duplicative Entry

Limited Visibility

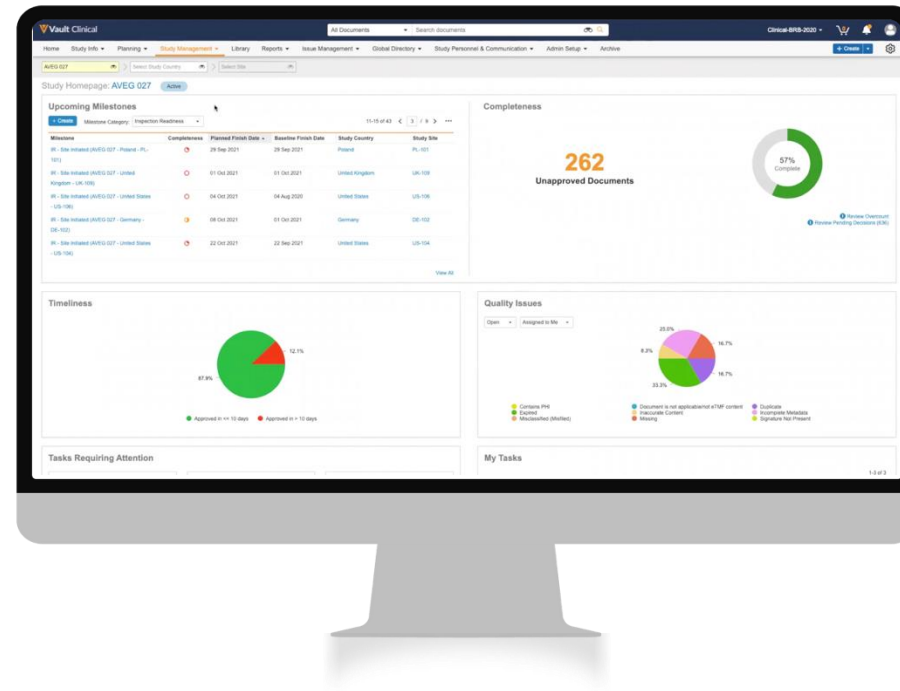
Clinical Operations

Standard platform for trial execution and site engagement



Veeva eTMF

Enable real-time inspection readiness, visibility, and control



Simplified
TMF Planning



Streamlined
TMF Activities



Relevant
Insights



Information
Sharing



Foundation
for Future

Veeva CTMS

Reduce complexity and accelerate trial execution



Streamlined
Trial Conduct



Simplified
Site Monitoring



Effective
Oversight



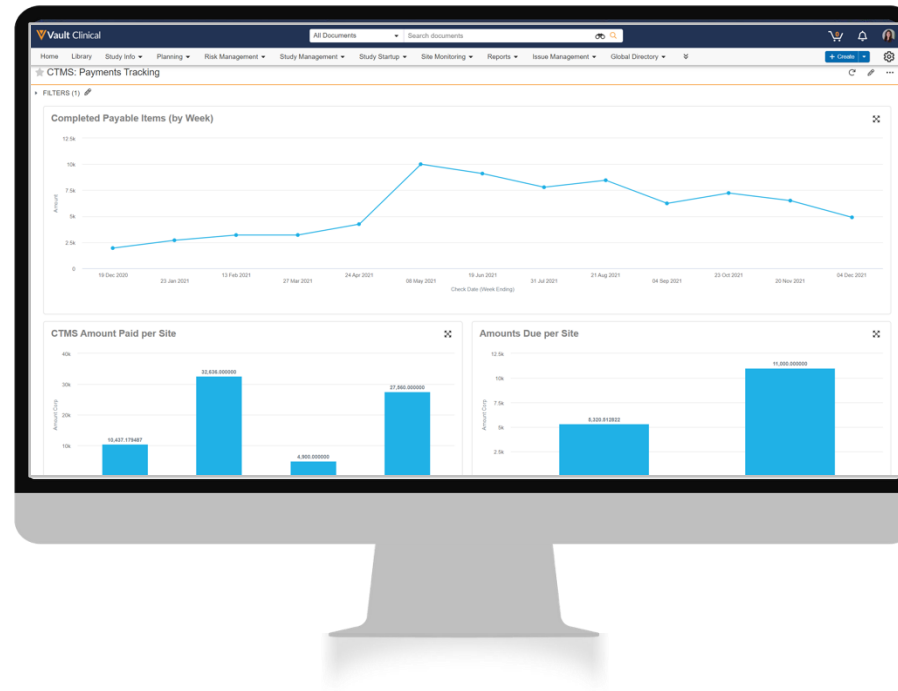
Optimized
Study Planning



Relevant
Insights

Veeva Payments

Pay sites faster and provide visibility to all study partners



Configurable
Fee Templates



Triggered
Payments



Robust
Workflows



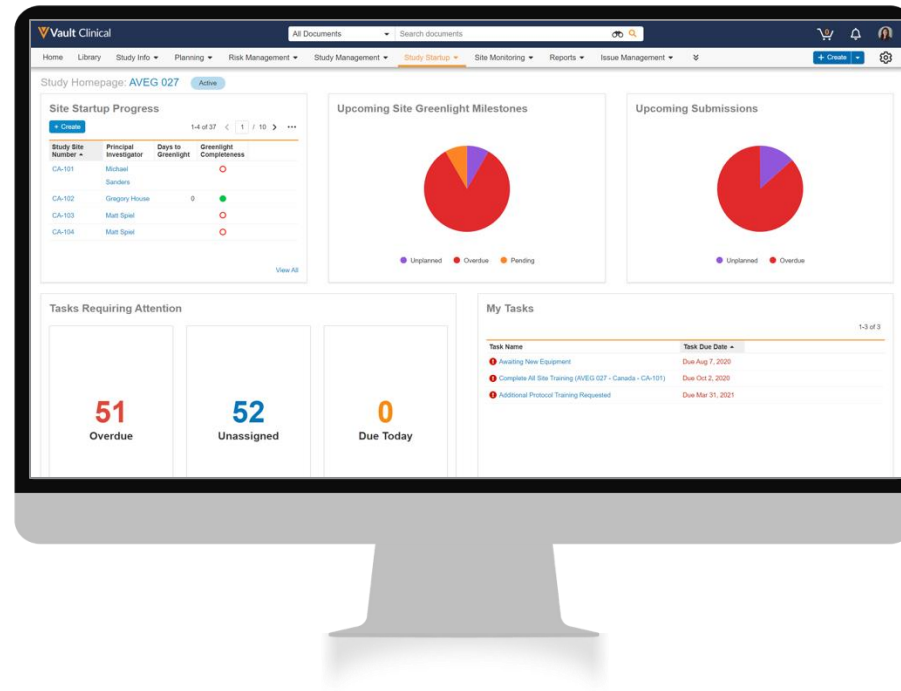
Relevant
Insights



Unified with
Veeva CTMS

Veeva Study Startup

Accelerate time to site activation



Automated
Feasibility Surveys



Intelligent
Country Workflows



Collaborative
Documents



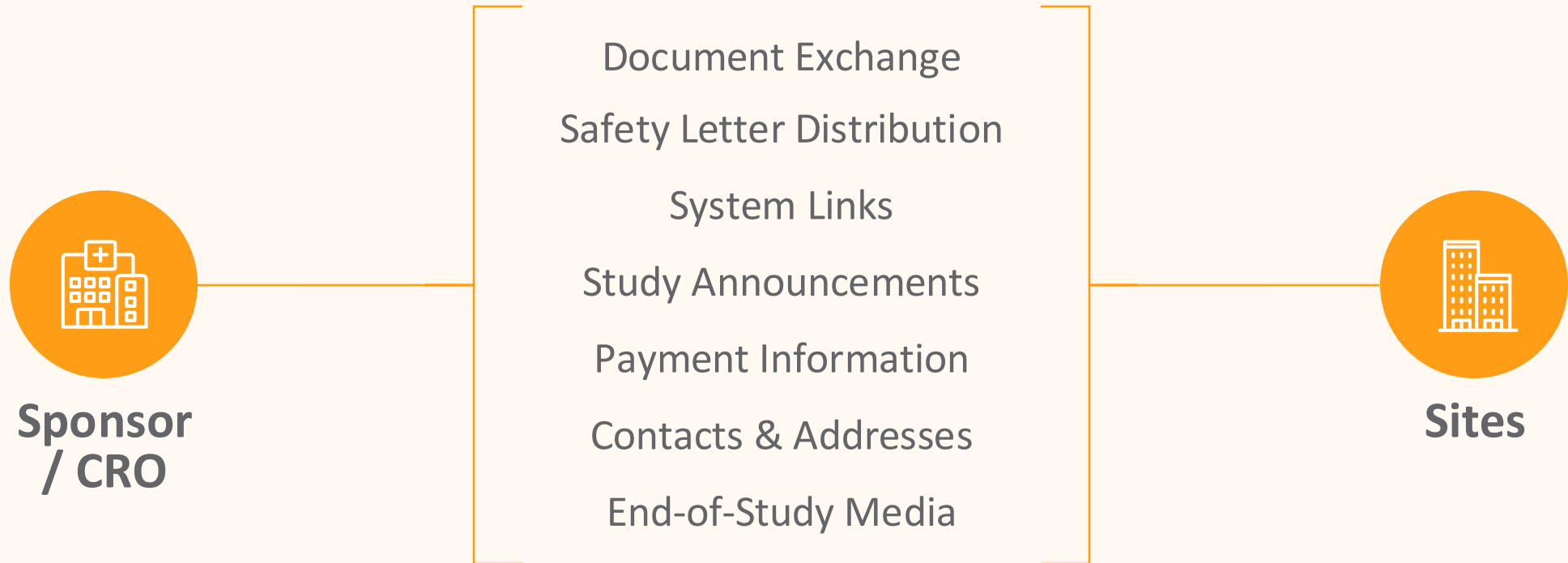
Streamlined
Negotiations



Relevant
Insights

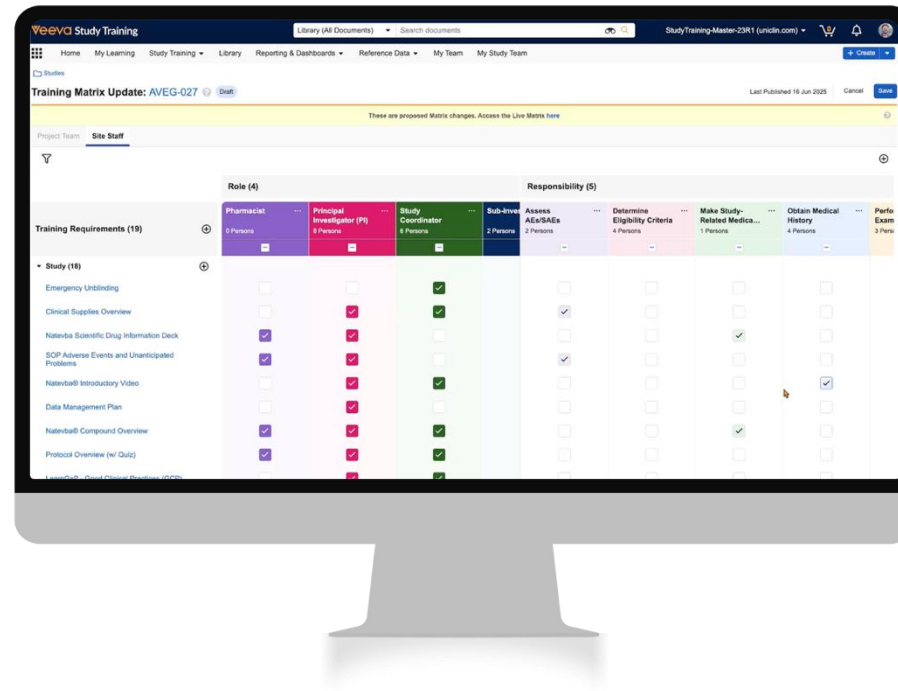
Veeva Site Connect

Simplify and standardize sponsor-site collaboration



Veeva Study Training

Streamline and automate study training



One Connected Solution



Streamlined Training Mgmt.



Visibility and Insights

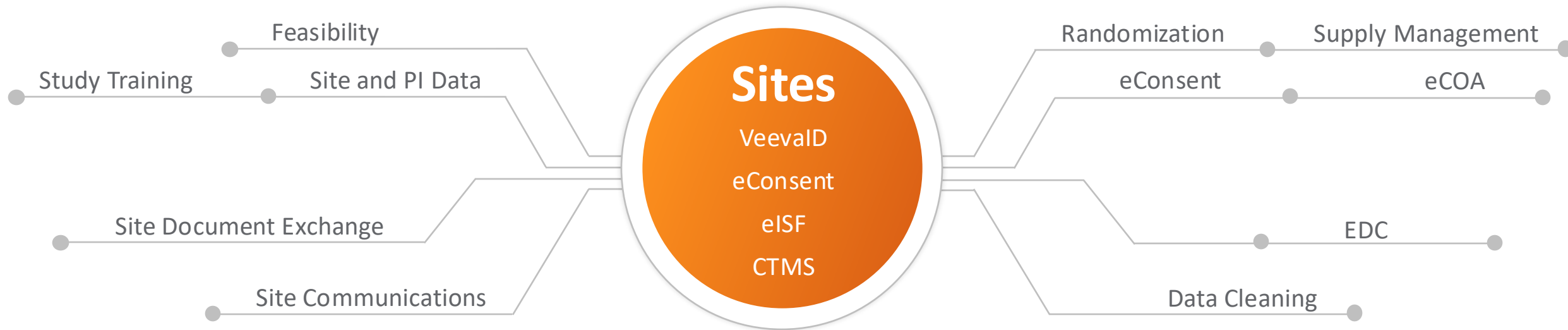


Training Reuse and Flexibility



Streamlined Reconciliation

Site Success Is Central to the Veeva Clinical Platform



CONSISTENT



CONNECTED



COMPREHENSIVE

How Sponsors Use Veeva to Reduce Site Burden

START-UP		CONDUCT		CLOSEOUT
Feasibility Administer industry-standard feasibility questions to create site and PI profiles	Doc Exchange Prepare document packages, flag overdue documents, and specify site actions needed	Training Assign training before site visits and track on-site activities	Safety Letters Distribute safety letters and track receipt and acknowledgement	EOS Media Transfer end-of-study media, including completed case report forms (CRFs)
—	—	—	—	—
Reduce asks of a site	Reduce site email volume	Improve the value of on-site time	Streamline site compliance	Simplify study closeout
▼ Study Startup	▼ Site Connect	▼ Study Training	▼ Site Connect	▼ Site Connect





DEMO



Thank you

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