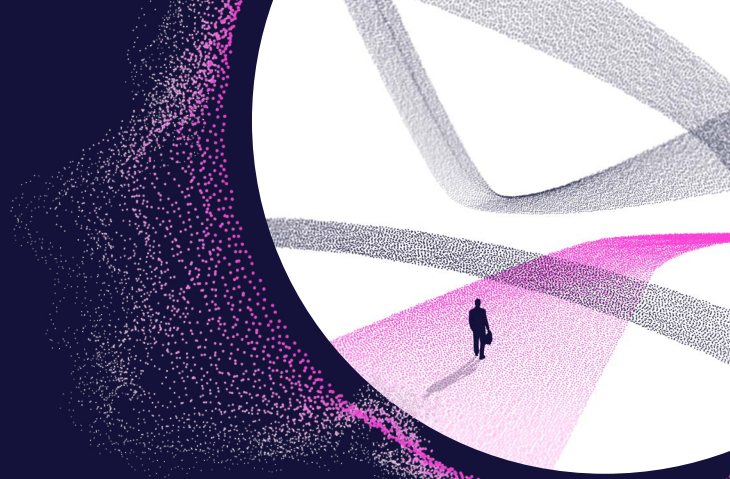
TRIALSCOPE DISCLOSE
CITELINE REGULATORY



Accelerated clinical trial disclosure starts here

With TrialScope Disclose, simplify, streamline and optimize the entire clinical trial disclosure process from initial registration to final results — mitigating risk, maximizing transparency, and ultimately moving healthcare forward.

Meet the most widely trusted solution for clinical trial disclosure management.

TrialScope Disclose — NOW INCLUDING Core Data!

Core Data is a unified registry submission platform using centralized content and all required global data fields to streamline the clinical trial disclosure process.

Benefits:

- Save time
- Enter and approve all data in one single place
- Keep data consistent
- Reduce questions from stakeholders
- Avoid duplications

- **37,000+** users managing **101,000+** trials on our platform
- Trusted by **16 of the top 20 industry** clinical trial sponsors
- Supporting customers responsible for **40%+** of industry studies registered on ClinicalTrials.gov, EudraCT, and CTIS
- Unrivaled experts in disclosure with **20 years of experience** dating back to the beginning of modern disclosure and transparency requirements

Optimize

Know exactly what, when and where to disclose

An intelligent disclosure rules engine automatically alerts you of impending deadlines, for compliance confidence and peace of mind.

Streamline

Simplify the entire process, from registrations to results

Centralize your trial data and use smart, customizable workflows for more efficient and accurate data entry, review & approval cycles, and registry submissions.

Simplify

Take the complexity and chaos out of compliance

Data-driven dashboards give you the snapshots you need for timely submissions and stress-free audits.

TRANSFORMING THE WAY YOU SEE THE WORLD

CITELINE.COM

Single Source of Truth

Unified registry submission platform for validated data collection and review.

Register the Right Data, in the Right Place, at the Right Time.

IMPROVE SPEED & ACCURACY
Automate submissions to ClinicalTrials.gov using validated fields

WORK MORE EFFICIENTLY
Slash data entry, review & approval times by up to 85%

MEET DEADLINES WITH EASE
Keep track of due dates with our proprietary disclosure rules library

TIGHTEN UP YOUR TEAM
Clarify tasks & timelines with role assignments & configurable workflows

COLLABORATE WITH CONFIDENCE
Work whenever, wherever with field-level commenting & verified e-signatures

REPORT YOUR WAY
Choose from predefined templates or ad-hoc creation

ALWAYS BE AUDIT-READY
You're well prepared with our full log conforming to 21 CFR Part II

REPURPOSE TRIAL DATA
Sync your data across forms and registries

A Scalable Solution for your Disclosure Needs

STANDARD FEATURES

- Auto-compliance Rules Engine
- Disclosure Workflow for CTgov and EudraCT
- Auto-submission to CTgov and Download to EudraCT
- Plain Language Summary and Study Synopsis Workflow
- TrialScope E-Learning
- Reporting Dashboards
- Upload Connector: CTMS and Adverse Events
- Unified registry submission platform

CONFIGURED FEATURES

- Unique Corporate Policy and Rules
- Custom Workflows
- Source System Integration
- Global Registries

Discover how to gain control of your disclosure processes, increase transparency and build your organization's reputation, starting with TrialScope Disclose.

To learn more about the advantages we can deliver to your company, please visit:
Citeline.com or email: **info@Citeline.com**