

# Prepare for **New Clinical Trial Disclosure Regulations** with TrialScope Intelligence

There is an increased spotlight on non-compliance of clinical disclosure regulations around the world. Regulatory bodies are raising the stakes for clinical trial sponsors of all sizes through new requirements and harsher penalties. Requiring forward-thinking and a regulatory knowledge base to successfully manage clinical trial disclosures.

**Eliminate the manual burden of hunting for clinical trial disclosure regulations and industry insights**

Our experts monitor, collect, curate, and analyze all new and updated requirements for you. Spend less time looking for and deciphering disclosure information—and more time on strategic, high-level tasks.

TrialScope Intelligence provides you with true global coverage, monitoring countries with and without current disclosure requirements. English translations and expert analysis create additional context and further your understanding of regulatory expectations.

## Key Content

Global — All in one place			
Country and registry requirements	Sources of truth	EN-language translations	Disclose Perspectives Analysis



TrialScope Intelligence also alerts you of any changes and conveniently consolidates them in one place for easy access. In addition, you can use action-oriented features to capture your trains of thought, communicate with your team and stakeholders, plan and strategize.

## Actions

Collaborate — With team members and experts alike	
Add Private Notes	Add Comments
Toggle Alerts	Reach SMEs Ask the Analyst



## Start with strategy. End with trust.



**Save time and resources:** Regain hours previously spent mining disclosure regulations for pertinent information. The TrialScope Intelligence experts curate disclosure requirements for you minimizing monitoring and manual work on your end.



**Mitigate non-compliance risk:** The consequences of failing to comply with disclosure requirements grow greater by the day, from heftier fines to longer jail times. By keeping up with ongoing regulatory changes, you can avoid damaging repercussions to your business and make any necessary adjustments to your SOPs ahead of time.



**Fortify public confidence:** Clinical trial participants want trial sponsors they can trust—and investors want honest, credible partners. Stakeholders and watchdog groups will appreciate a more robust, transparent, and compliant clinical trial disclosure process.



**Enhance collaboration:** The disclosure process is often disjointed, making collaboration across time zones and workspaces unnecessarily challenging. With TrialScope Intelligence, all colleagues can stay up to date on compliance changes by setting alerts and tagging team members in the comments.

Ready to reinforce new habits  
and build stronger clinical trial  
disclosure processes?

**Get your free demo today.**

To learn more about the advantages we can deliver to your company, please visit:  
[Citeline.com](https://Citeline.com) or email: [info@Citeline.com](mailto:info@Citeline.com)

