

Use Case

# Disclosure Transparency Across Local and Global Registries

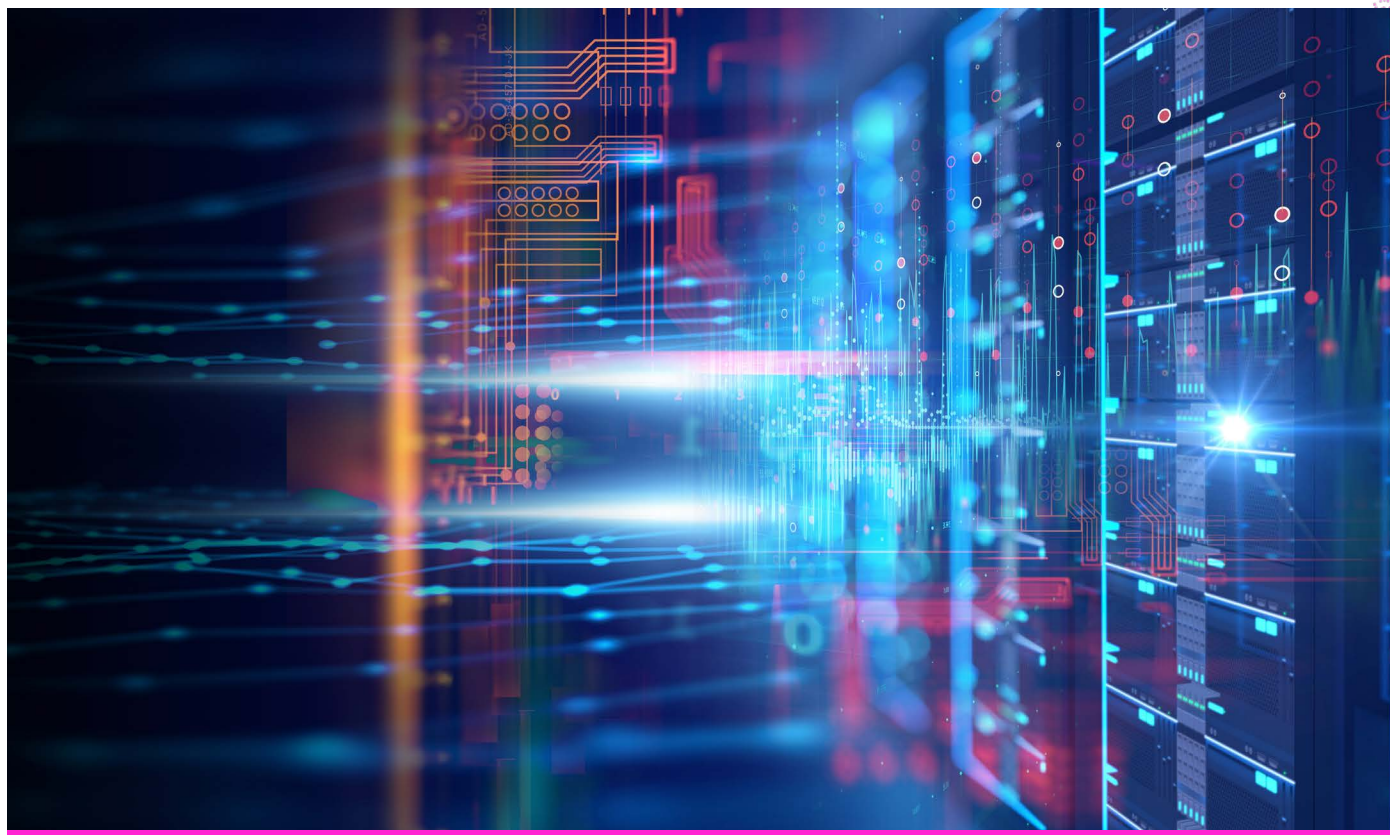


## Use Case | Disclosure Transparency Across Local and Global Registries

### The Need

Clinical trial disclosures on local registries are often managed at the local level; the process is rarely transparent from a global perspective. Sponsors need to reduce the risks of information being accidentally disclosed on local registries by maintaining better central control of disclosures.

A clinical disclosure team leader developed a project to improve alignment of disclosure filings across registries and countries. The first step of the project was to swiftly get a full understanding of all mandatory disclosure requirements worldwide.





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### The Solution

**TrialScope Intelligence**, with its global coverage of disclosure requirements and 59 registries across 193 countries, provided the critical knowledge needed for the first step to this project. The client could customize TrialScope Intelligence's centralized, interactive repository of requirements to focus specifically on the locations relevant to the client's company.

Using TrialScope Intelligence removed the burden of manually gathering accurate and up-to-date requirements, giving the company back time to focus on its strategy for planning timely and compliant clinical trial disclosures and reducing the risks of accidental local disclosures.





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### The Benefits

Ultimately, TrialScope Intelligence allowed the clinical disclosure team member to:



**Save time and resources** by using a centralized system to assess mandatory regulations in each country and investigate each registry required globally



**Improve transparency** across registry filings by seamlessly comparing requirements for each relevant registry so they disclose only as necessary information



**Mitigate noncompliance risk**, avoiding fines and damages to the company's reputation by ensuring each registration contains precisely the data required by local and global disclosure regulations





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