

Enabling Pharmaceutical Companies to Automate Data De-Identification for Secondary Use



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The Cost of Transparency

RESOURCES NEEDED





Time

Reacting to new regulations and developing new processes takes time.



Money

Both outsourced and internal methods require additional costs.



Knowledge

Keeping up-to-date with current regulations can feel like a full-time job.



Public Education

Transparency enables the public to make informed decisions regarding their health care.



Trust

Being transparent with trial data builds trust in the clinical trial process.

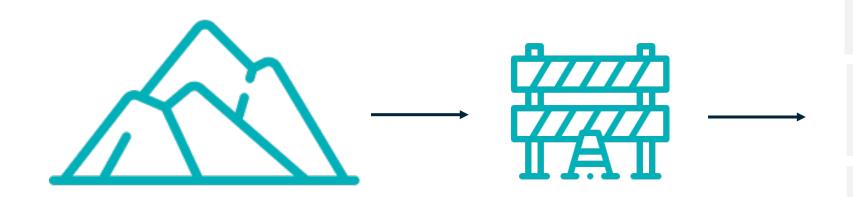


Secondary Research

Empower medical breakthroughs by utilizing existing data to minimize duplicative efforts and reduce timelines.



Thinking Beyond Regulations



Completed Study Data

Compliance Standards
Roadblocks

Exploratory analysis to support future studies

Minimize duplicate efforts

Medical Breakthroughs

Potential



Automating De-Identification at a Large Scale | Case Study

A global sponsor enlisted d-wise to provide a proof-of-concept capability to **accelerate the flow of data to and from an internal data lake** through d-wise's Blur anonymization solution, providing sufficiently anonymized content to achieve re-identification risk levels appropriate for internal secondary use.

Challenges

What to Anonymize
Internal Process Alignment
Scalability
Risk Threshold Strategy

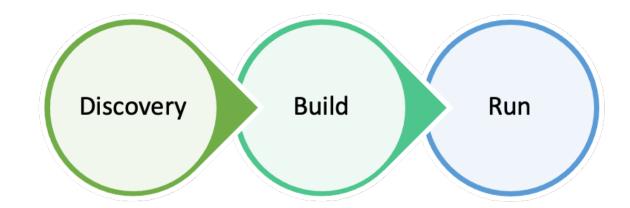
Solution

Automated de-identification of datasets through Blur



d-wise Approach

- Discovery, Build, Run Process
- Master Template Format
- Audit Trail
- Variable Analysis
- Verbatim Text
- Inbound Folder Structure for System Input
- Outbound Folder to Receive Final, Resorted Data





Run Phase

Manual

 Place study folders into Inbound folder

Automated

- Template applied
- Calculating Match Score
- Anonymization Report
- Resorting datasets
- Exporting into study folders in Outbound folder

Results

84

Studies Anonymized 30

Minutes

Manual Task



Contact

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