

Cerevel Bridges Gap Between Regulatory and Clinical



Business Challenges

- Disconnected data management and entry between clinical and regulatory
- Overlapping documents required manual CrossLinks & duplicate uploads
- Resulted in availability delays, misaligned data, and greater risk for human error



Veeva Solution

- Vault Clinical Operations to RIM Connection will facilitate document delivery of 1572s and investigator CVs from CROs to regulatory
- Study-related documents (e.g. protocols and IBs) now populated in real-time for inspection-ready state
- Minimal training required



Tangible Benefits

- Product and study data are aligned with a clear data flow
- Users are saving time and maintaining compliance with real-time availability of key study documents and a single source of truth
- Regulatory and clinical Vault administrators are more engaged



We have just scratched the surface on the number of interlinked clinical and regulatory processes that can be enhanced with this connection.



Dee DeOliveira, **Director of Global Regulatory Operations at Cerevel**

