

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**

