

Veeva R&D and Quality

# SUMMIT ONLINE

Join us online, October 13-14, to hear market trends in the industry, see innovative technology, and learn from life sciences experts.

Register Now

## Top 3 Reasons You Should Join Us Online

Learn best practices from customer case studies


+

Get a sneak-peek into innovative products with live demos

+

Join discussions with peers and industry experts

### Access to

  
**80+**  
sessions

  
**50+**  
customer speakers

  
**3,000+**  
life sciences professionals

### Keynote



- Amy Abernethy, principal deputy commissioner from the FDA, joins as the keynote speaker to discuss speeding drug development through technology innovation.

## Highlights From Each of Our Six Tailored Tracks

### Clinical Operations

- Results from Unifying Clinical Operations, AstraZeneca
- Real-World Audit Results, Seattle Genetics

### Clinical Data

- Global Roll-out of CDMS, Eli Lilly
- Enabling Data Managers to Build Studies, Alcon

### Quality

- Quality Transformation, Bristol Myers Squibb
- Global Document Management, GlaxoSmithKline

### Regulatory

- Regulatory Transformation, Amgen and Sanofi
- Vault Submissions Publishing, Myovant, Viela Bio, and Dermavant

### Vault Platform

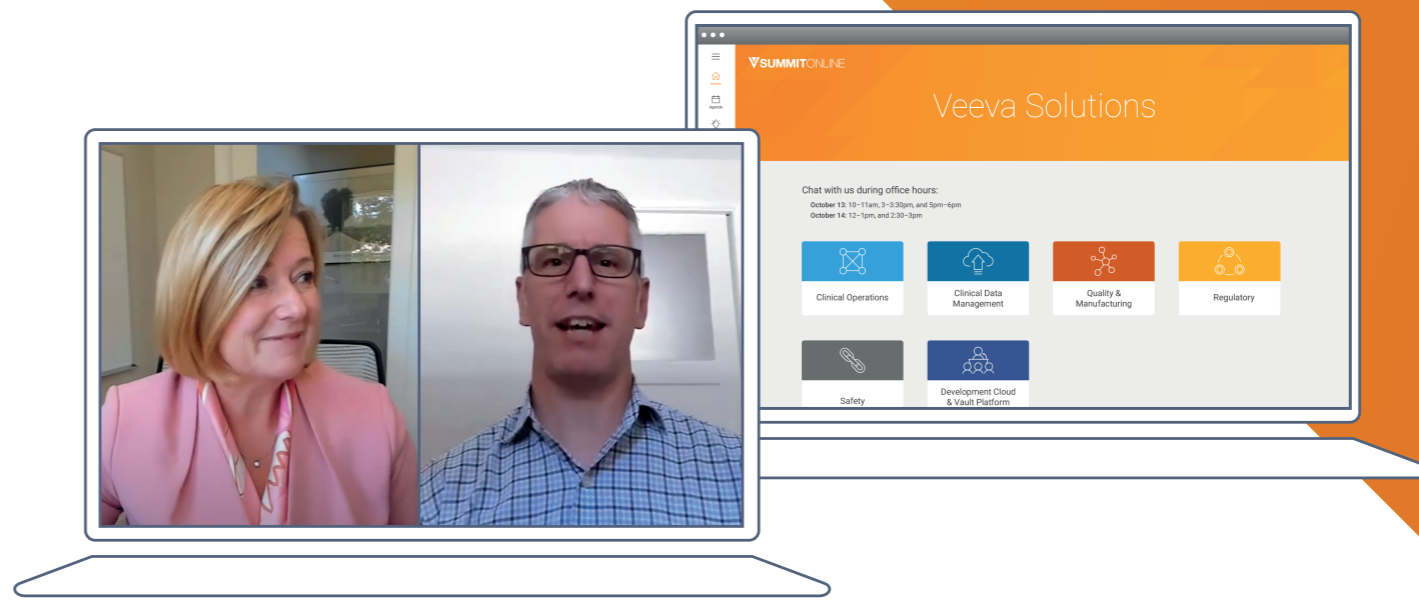
- Connecting Vaults across the Development Cloud
- Vault Platform Roadmap and Vision

### Safety

- Veeva Vault Safety Suite Overview and Vision
- Case Management When Outsourcing, G1 Therapeutics



Ask questions via live chat and interact with product experts.



## What EU R&D Summit Attendees Said

Summit brought together real users and was extremely interactive. Plus I picked up some great ideas.

I leave Summit upbeat and excited about opportunities to improve operations going forwards.

Virtual Summit exceeded my expectations. The excellent presentations by industry leaders sparked ideas to take back to my own organization.

Register now  
[veeva.com/rd-summit](http://veeva.com/rd-summit)