



5,000+



350+

Clinical data, simply

Trial management solutions

Seamless collaboration
meets actionable data



cubeCTMS

Your trial oversight hub

- Detect signals early
- Micro-tracking and macro-reporting

cubeTMF

Be inspection-ready

- Full trial documentation control
- Site master file management

cubeRBQM

Real-time insights

- Monitor risks proactively
- Greater focus with interactive dashboards

cubeLMS

Train with confidence

- Create tailored training courses
- Track compliance effortlessly

cubeCDMS

The cornerstone of
clinical data innovation



Streamlined data management

- Detect deviations automatically
- Native risk-based monitoring

Enhanced data quality

- Minimize queries, lighten workload
- AI-powered medical coding

Simplified self-service setup

- Library with 55+ pre-built eCRFs
- No programming required

Unified CRScube platform

- No more data entry duplication
- No need for data reconciliation

Data capture solutions

Streamlined data collection and patient management



cube **IWRS**

One-click randomization

- Flexible inventory management
- Accelerated RTSM setup

cube **DDC**

Paperless precision

- Timely, accurate data entry
- Portable and offline mode enabled

cube **PRO**

Patient insights, simplified

- View patient data in real-time
- User-friendly mobile app

cube **CONSENT**

Positive patient engagement

- Easy informed consent process
- Interactive content platform

cube**SAFETY**

Ensuring vigilance through data



Submit with confidence

- Prevent duplicate case entries
- Safety signals from various sources

Meet regulatory standards

- FDA, EMA, CDE, MFDS and PMDA
- ICH E2B and regional guidelines

Track with precision

- Instant access to agency feedback
- Real-time submission status

Boost efficiency

- Automated data entry
- Easy data import and export



Empowering clinical innovation,
one trial at a time



Experience
CRScube

www.crscube.io
marketing@crscube.io

