

# Speeding EDC build with rapid study design

Imagine designing, approving, and building **full** EDC studies faster than ever with CRF Creator from Pinnacle 21.

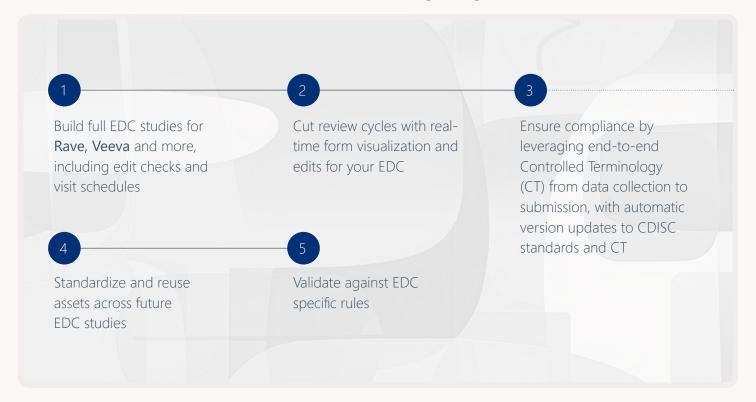
The CRF Creator platform gives data managers the tools to create compliant, standardized CRFs, empowering clinical teams to start studies sooner.

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|-------------------------------|--|
| Pinnade 21° Adverse Event     | Additional Metadata   Annotations   Show Edit Checks/Derivations |
| Subject Name<br>Adverse Event |  |
| Date of visit:                |  |
| Adverse Event description     |  |
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Standardization and reuse Multi-EDC flexibility

Real-time form visualizations One-click EDC build

# 5 Reasons to use CRF Creator for designing EDC studies



#### Core features

- Design forms with:
  - Question positioning
  - Form dynamics (e.g. radial drop-downs)
  - Nested questions
- Preview forms for your EDC
- Deploy a fully functional study with:
  - EDC specific edit checks
  - Visit schedules
- Manage Controlled Terminology (CT) across your EDC, SDTM & Adam
- Auto-update changes to CDISC standards & CT
- Automate annotated CRFs and edit-preview changes in real-time
- Standardize assets for reuse, including:
  - Forms
  - Visits & visit schedules
  - Edit checks
  - Controlled Terminology (CT)
  - Annotations

## **EDC** integrations

















### Business benefits of Pinnacle 21 CRF Creator



50% faster EDC build: Speed database build with accelerated review cycles and reuse of standardized assets.



relevant EDCs for different study phases.



Save time and costs: Reduce resource-intensive processes and repetitive manual work through automation.



Improve quality and consistency: Enhance quality across studies and reduce risk with standardization and reuse.



Ensure compliance: Stay aligned with CDISC standards and automate Controlled Terminology updates.

Ensure faster startups, higher quality, and accelerated study builds - in one centralized platform.

Learn more about Pinnacle 21 CRF Creator

#### **About Certara**

Certara accelerates medicines using proprietary biosimulation software, technology and services to transform traditional drug discovery and development. Its clients include more than 2,400 biopharmaceutical companies, academic institutions and regulatory agencies across 70 countries.