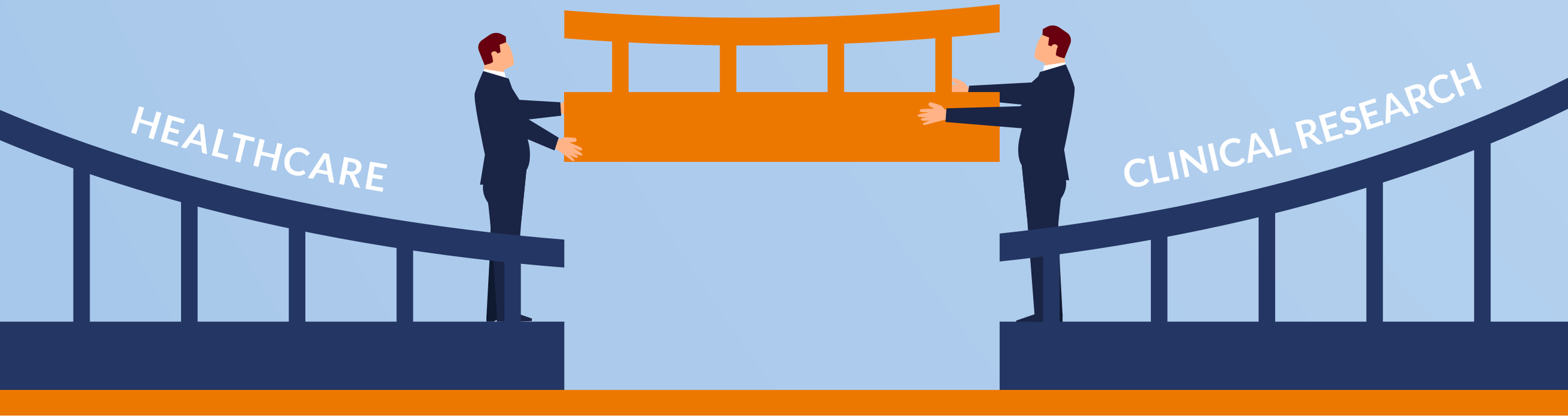
Introducing OpenClinica Unite™

Bridging the gap between healthcare and clinical research



www.openclinica.com

OpenClinica

Manually transcribing patient data is extremely inefficient, time consuming, and highly error prone.

Running a clinical trial requires lots of data that already exists in your EHR. This critically important data serves as the source record for clinical trial participants, but must be manually transcribed into the study database. Finally, there is a way to streamline this process.

Introducing OpenClinica Unite™
A powerful and easy way to leverage EHR data for your next study.

Imagine a world where, with a single click, you could get EHR data into your study database. Because now you can.

Eliminate high-risk manual data entry

Reduce monitoring cost and effort

Get faster access to data

Research sites can:

- Save countless hours expended by overburdened coordinators and clinicians — reduce routine tasks from hours to seconds!
- Enhance patient safety and care by tracking important metrics
- Leverage **Unite™** for multiple trials and sponsors
- Participate in research more easily and profitably

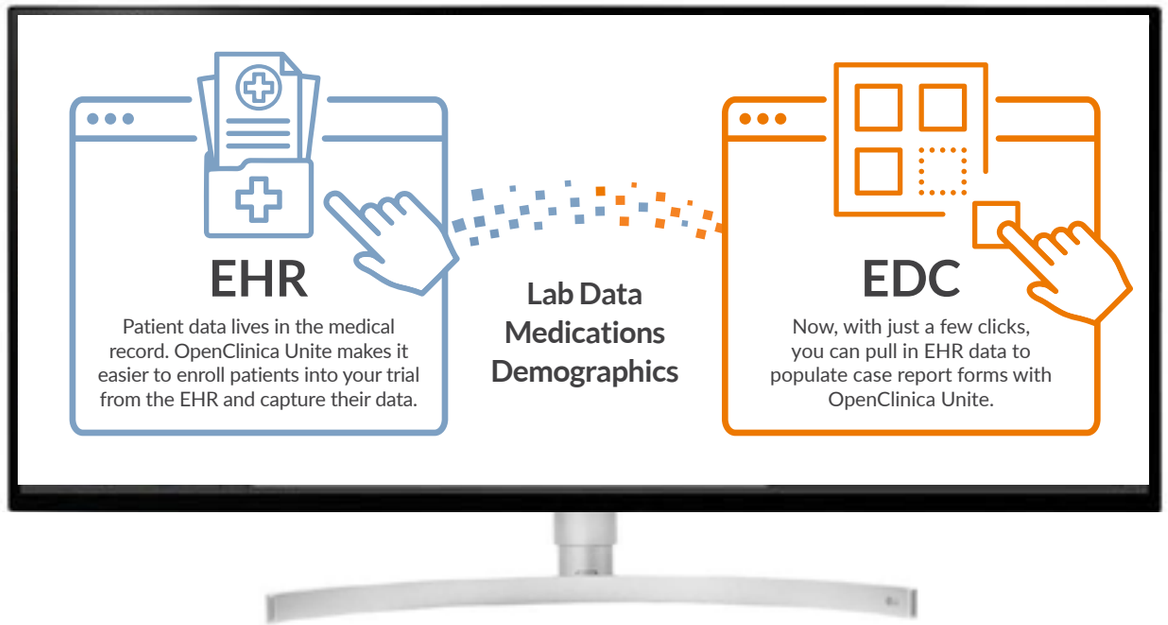
Sponsors can:

- Obtain higher quality data that is cleaner and more complete
- Get faster access to study data
- Reduce the costly practice of 100% SDV
- Increase site satisfaction and compliance

OpenClinica Unite™ is a seamless solution that can be directly tied into your existing EDC/study database infrastructure.

Meet OpenClinica Unite™

OpenClinica Unite™ expands the use of electronic health record data in clinical research and greatly reduces the burden of data entry for sites. With OpenClinica Unite, you can securely retrieve data from your EHR to automatically populate trial case report forms.



Seamless user experience

- Single sign-on with EHR - clinicians, investigators, and coordinators can launch the eCRF from inside the patient chart within the EHR
- Available in both desktop and mobile to allow users to safely, accurately, and efficiently work with research data in the way that makes sense for them
- EHR patient - EDC participant linkage automatically maintained
- Real time data visualization for use at the point of care (e.g., track key metrics)
- Puts EHR data, traditional eCRF, eCOA, and ePRO data in one place at users' fingertips

Rigorous compliance

- Compliant with 21 CFR Part 11 standards and ICH GCP
- Compliant with HIPAA and GDPR
- Traceability to the source (data originator, data element identifiers per FDA 2018 EHR guidance)
- Comprehensive information security program
- Compliance and security artifacts available for customer review

Easy to implement

- Secure, standards-based technology accepted by health system IT departments and backed by major EHR vendors
- User-driven access to data (easily auditable on source side)
- Can be implemented in one day and used with any EDC
- Reuse implementation across multiple trials and sponsors
- Supports hybrid trials - not all sites need to use it

Highly customizable

- Visualize your data with embedded OpenClinica Insight reporting that allows clinicians to view the participant record at a glance and monitor key trends to enhance safety, data quality, and support optimal clinical care
- Unite your data with highly configurable design options that map data elements such as lab results, medications, and demographics to ontologies in your EHR (i.e., LOINC, RxNorm, SNOMED, etc.)

Curious to learn more?

See how OpenClinica Unite™ is transforming clinical research for a high-profile COVID-19 study. We'll show you a real use case and discuss how Unite can benefit your clinical research based on your specific needs.

STOP WASTING PRECIOUS TIME AND RESOURCES!
Get back to doing what you do best: improving lives through innovative, transformative research.

Request Free Demo