

E360™ CLINICAL DEVELOPMENT: REAL-WORLD INSIGHTS TO INFORM PROTOCOL DESIGN

Discover the world’s largest healthcare research platform¹...

E360™ Clinical Development is the industry leading SaaS technology platform offering best in class real-world insights to inform protocol design and state-of-the-art analytical tools for effective trial optimization.

CURATED HEALTHCARE DATA SOURCES

- Over **500 million** patient lives across eight countries
- Seamless multi-country analysis/studies execution: OMOP Common Data Model (available on request) enables to **build your analysis/study once and deploy anywhere**

EXAMPLES DATASETS AVAILABLE

| COUNTRY | TYPE |
|----------------|---|
| AUSTRALIA | • Ambulatory EHR |
| BELGIUM | • Ambulatory EHR |
| CANADA | • Ambulatory EHR |
| FRANCE | • Ambulatory & Specialty EHR |
| GERMANY | • Ambulatory & Specialty EHR |
| ITALY | • Ambulatory EHR |
| UNITED KINGDOM | • IQVIA Ambulatory EHR |
| UNITED STATES | <ul style="list-style-type: none"> • Ambulatory EHR • P+ Health Claims • Oncology EHR • Open Source Claims • CMS Synpuf Claims |

SPEED MATTERS

E360™ Clinical Development enables instant global feasibility on large scale EHR.

COMBINE SPEED, TRANSPARENCY AND ANALYTICAL POWER IN ORDER TO:

- Access Ambulatory and Specialty EHR seamlessly across US, Canada and European countries
- Simulate complex protocols, end-points and relative time frames
- Establish detail country feasibility, local treatment and patient visit patterns
- Conduct precise “what-if” analysis and receive feedback in real time

RESULTS MATTER

CLIENT #1

- Protocol optimized against real patients’ data in hours
- Identified operational risks due to differences in real-world clinical practice in the early planning phase of the trial
- Developed the right strategy to mitigate operational risks before recruitment
- Avoided potential delays and costly amendments during recruitment

CLIENT #2

- Verified protocol design against real patients’ data in hours
- Confirmed protocol feasibility and patients’ availability across countries in real time

¹ In terms of the number of de-identified patient records contained

USING ELECTRONIC HEALTH RECORDS AND E360™ ACROSS PROTOCOL DESIGN AND TRIAL DESIGN

TPP/CDP²

- Understand the burden of disease
- Identify all local comparators
- Characterize patient populations and geographic variations
- Refine the overall market potential for particular TPP

PROTOCOL FEASIBILITY

- Qualify the exact available population
- Simulate exact impact of specific I/E criteria
- Test complex protocols including relative time frames

INITIAL STUDY DESIGN

- Simulate recruitment rates based on real patient visits
- Understand impact of seasonal variation
- Characterize sites and available patients

FINAL DESIGN TO START-UP

- *Input into CTOS Design for protocol design and development*
- *Leverage CTOS SiteOptimizer for site and investigator selection*

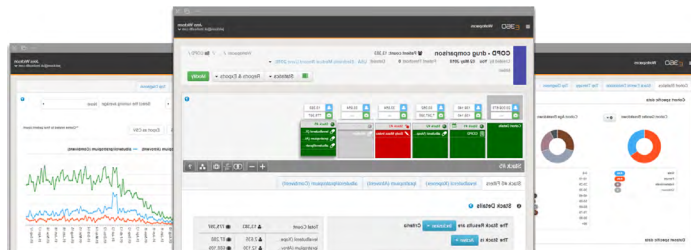
EARLIER USE OF A DATA-DRIVEN APPROACH DRIVES IMPROVED SUCCESS RATE

² Target Product Profile/Clinical Development Plan

EXAMPLES OF CAPABILITIES AVAILABLE

PROTOCOL FEASIBILITY

- Profile and find your next real-world data assets
- Interrogate any element of EHR data (diagnosis, drugs, labs, etc.)
- Define complex protocols re-using existing codelists and disease definitions
- Simulate even the most complex criteria including relative time windows
- Instantly test across multiple countries for global feasibility



ADVANCED REPORTING

- **Geo Visualization**
Identification of geographically favourable sites for clinical trials
- **Incidence and Prevalence**
Report disease incidence and prevalence on specific cohorts
- **Attrition**
Visualize the patients funnel for a specific cohort
- **Patient Timeline**
Visualize the patient's journey and when they become eligible and recruitable in their own healthcare journey
- **Data Completeness**
Interpret EHR properly and understand how well key clinical variables are captured in EHR

