

# Pharmaceutical Companies Accelerate Clinical Trial Information Flow



## Company

**US Pharmaceutical Companies**

## Industry

**Life Sciences**

## Application

**Clinical Trials**

*"Sites like it. The reliability and quality are impressive. We have more data, faster."*

Eric van Son, CTO

## Challenge

- Takes weeks to collect and process clinical trial data
- Majority of data is collected on paper and must be re-keyed adding to time and errors
- Visibility of results is critical throughout the trial enables adaptive trial process
- Delays can result in losing "first-mover" advantage
- Inaccuracies can result in delays, health risks or losing the value of a new drug
- Other data capture solution require site staff to become data entry clerks

## Solution

- ExpeData Digital Writing Platform
- 21 CFR Part 11 functionality
- Closed and secure system
- Searchable and secure documents
- All data is encryption for needed security
- Paper source document (CRF) is retained for backup

## Results

- Increased site productivity, monitor's cost and time reduced by 30%
- Near real-time data exchange between sites and centralized coding, reduces time to deliver data to sponsors
- Handwritten data is available in seconds
- Trials close sooner providing faster results and faster approvals
- Easy to deploy, easy to use, high adoption rate and the "Sites like it"