

# PUT AN END TO eCOA MISCONCEPTIONS



## Regulatory bodies support eCOA data

Regulatory bodies have now made their stance clear toward eCOA solutions:

- **They recommend** the use of electronic solutions to collect clinical data endpoints
- **The FDA has confirmed that** the electronic capture of clinical source information is preferred to a paper-based system

# 5



## A successful transition to digital technologies

Not all sites and patients are equal when it comes to technology:

- **Customized education and training solutions** should be provided
- Patients and sites should be reassured and supported with **adapted level of advice** and **clear documentation**

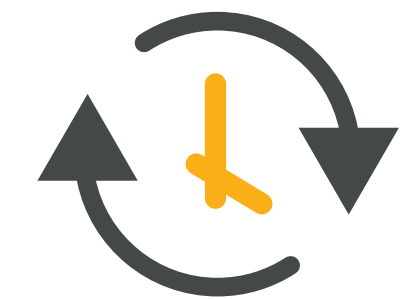


## eCOA implementation: a valuable investment

eCOA use reduces the hidden costs compared to a paper-based system:

- **eCOA implementation reduces the costs** related to non-compliance, poor data quality, data management, site monitoring and study closure activities
- **More sensitive results with eCOA**, thanks to the reduction in error rate and the improvement of data quality

## reasons to adopt eCOA solutions



## eCOA solutions ensure time savings

eCOA implementation can allow weeks to be saved:

- eCOA tools make data **attributable, legible, contemporaneous, and accurate**
- **Robust project management** of the trial start-up activities means that **study timelines are met**



## eCOA optimizes data collection

eCOA ensures more data & better data integrity and accuracy compared to a paper-based system:

- Reduced risk of losing data thanks to **data synchronization**
- Efficient and appropriate **eCOA back-up plan** available

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