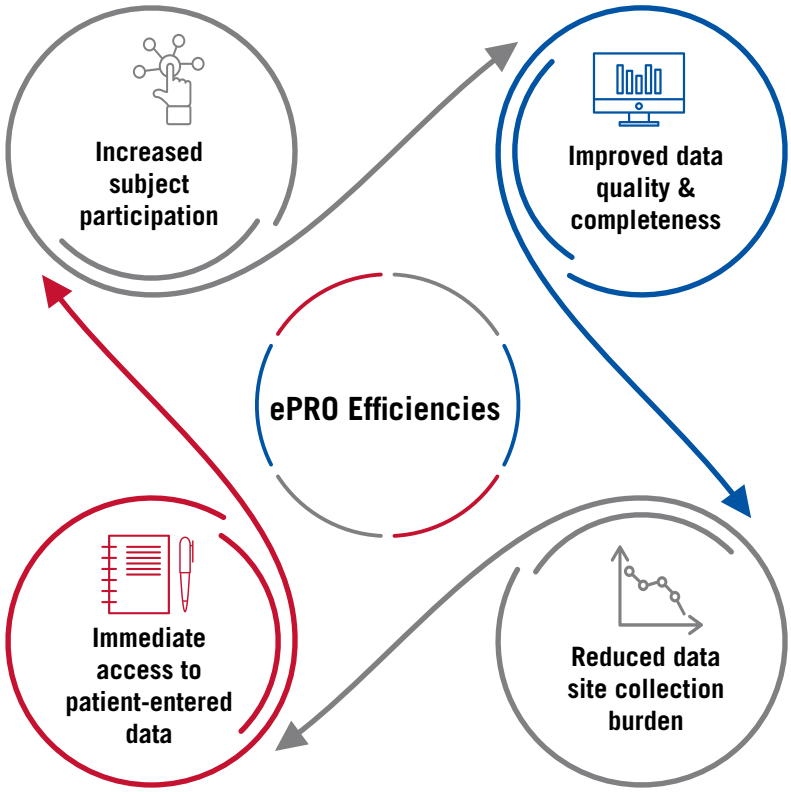
ePRO

Increase Patient Engagement and Help Ensure Compliance During Any Clinical Trial

Managing patient reported outcomes can be time consuming and costly. The *iMedNet* ePRO module streamlines the process by making onsite and offsite data captures easy for both Patients and Research Coordinators. ePRO allows subjects or coordinators to intuitively enter study-related information, and is accessible - anytime, anywhere.

Simplify subject participation and protocol adherence with:

- Direct subject sign-on
- Patient To Do Lists
- Automatic email notifications
- Secure data collection



Our Commitment

MedNet is committed to empowering life sciences organizations by providing the most agile, efficient, and effective EDC and eClinical technology in the marketplace.

Through the delivery of our robust suite of capabilities, we are proud to assist clients worldwide in enhancing and streamlining clinical trial processes and improving patient outcomes.

iMedNet™ eClinical

Comprehensive
A single unified platform

Flexible
Superior built-in config*urability*

Cloud-Based
High availability and scalability

Secure
HiTRUST CSF platform and Privacy Shield certified provider

Proven
Successfully deployed worldwide

Road Map

Future enhancements to *iMedNet*'s ePRO.

- Patient To Do List: Configurable end window
- No Email login: More accessible to patients without emails

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