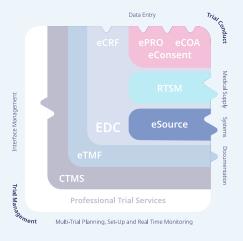
esource



ENHANCING CLINICAL DATA MANAGEMENT WITH OUR ADVANCED eSOURCE SOLUTION

Streamline your data collection and reduce costs with oomnia eSource

Our eSource solution has been crafted to simplify your clinical trial workflow. It enhances transparency, reduces manual workload, increases accuracy, decreases the need for manual on-site SDV, and fully enables remote monitoring. From regulatory compliant direct data capture and transfer to EDC and electronic health records, wearable technologies to real-world data and mobile applications, oomnia captures all electronic source data in real-time, eliminating the need for manual entry.

- ✓ Accurate and reliable data
- ✓ Integration with EDC/EHR, wearables, and more
- ✓ Flexible and scalable
- ✓ True remote monitoring

BENEFITS OF OUR UNIFIED eSOURCE

Instantly capture and manage clinical data



INCREASED ACCURACY

Increase data accuracy by eliminating costly transcriptions errors and outdated workflows. Users enter data into the eSource and it is automatically mapped to the eCRF or EDC and even to the EHR.



COST SAVINGS

Reduce costs by dramatically reducing SDV. With automatic data transfer from eSource to the oomnia EDC, data is always the same as the source. Our web-based oomnia eSource fully supports remote monitoring.



CONFIGURABLE

Users can customize data forms, permissions, and workflows to suit specific trial needs. Stakeholders benefit from secure, tailored portals with advanced role-based access controls that safeguard data integrity.



MUTLTIPLE DATA SOURCES

Transparently collect data from most anywhere. oomnia eSource enables real-time integration with wearables, hospital devices, LIMS, and more. Built on UMLS, HL7, FIHR, and LOINC standards, communication with any connected device is a breeze.



OFFLINE DATA COLLECTION

All the benefits of a web-based platform with offline capabilities. Generate data collection PDFs and send them to staff or patients to complete remotely. Upload them to complete eSource data collection instruments.



CONTEMPORANEOUS DATA

Decrease or eliminate the time from assessments to eCRF data entry. Data is visible and reportable from the time of entry into eSource and transferred to the eCRF. No waiting for staff to update eCRFs or the EDC.





ADVANCED FEATURES OF OOMNIA eSOURCE

Unlock the full potential of your clinical trials

INCREASED EFFICINECY AND REDUCED COSTS		
FEATURES	DESCRIPTION	
Streamline and accelerate data capture	Enter data into the eSource instead of paper CRF, and the data is automatically transferred to the eCRF or other EDC forms	
	Capture COAs from participant wearables or other connected devices directly in the eSource and automatically transfer to EDC	
	• Data is entered into the EDC	
Eliminate redundancy and reconciliation	Eliminate manual data entry into EDC and, if required into the EHR/HIS	
	No reconciliation required	
Reduce manual workload	No necessity to enter the same data twice	
	Data is entered only once into the eSource and transferred into the clinical database	
Maximum accuracy	Hard and soft data validation rules applied directly to the source	
	Eliminate manual transcription from paper CRF to eCRF and the errors that are be introduced	
Increased monitoring efficiency	Enable remote monitoring through a cloud-based system	
	Eliminate the need for SDV as data is transferred directly to the EDC, thus ensuring 100% accuracy and congruence with the source	
	Reduce queries	

FLEXIBILITY AND SCALABILITY		
FEATURES	DESCRIPTION	
Scalable data storage and processing capabilities	Handle growing data volume effectively, ensuring uninterrupted data processing, storage, and retrieval	
Offline eSource data capture	Complete data capture PDFs complete with mapping and data validation rules, save locally for upload to the eSource at a later date	
Adaptability to different trial designs and protocols	Flexible enough to accommodate different trial designs and protocols	

 Allow monitors to focus on the quality of the sites and not on rote SDV

EDC INTEGRATION	
FEATURES	DESCRIPTION
Immediate data transfer to EDC forms	 eSource to EDC Mapping using CDISC standards oomnia eSource solution is directly mapped and integrated with our oomnia unified clinical research software Data is transferred seamlessly to participant eCRF or any other EDC document
Data transfer to EHRs/HIS	 Transfer data from the eSource solution directly to Electronic Healthcare Records or Hospital Information System Built-in medical thesaurus Out-of-the-box HL7 and FIRH standard support for healthcare data transfer and mapping
Laboratory information management system (LIMS) integration	 Supports direct data transfer to eSource from LIMS to eSource for a single source of truth Out-of-the-box LOINC support both in the eSource as well as in the oomnia unified clinical research software

ACCURATE AND RELIABLE DATA COLLECTION		
FEATURES	DESCRIPTION	
Customizable data capture and forms	 Advanced and crucial ability to customize data capture methods and forms to suit specific trial requirements Utilize forms or questionnaires Direct data capture from IoT wearables Integrate with existing hospital devices for automatic data transfer 	
Real-time data access and reporting	 Data is immediately transferred from the eSource Improves management of clinical trial with real-time data access and analysis which allows for timely decision-making Data immediately available for export, graphical reports, or statistical analyses 	
Robust audit trails	 All actions by all users recorded in an event log Detailed audit trail exportable at any time for all data captured, whether in manually in forms or automatically from IoT enabled devices, or from other sources 	



