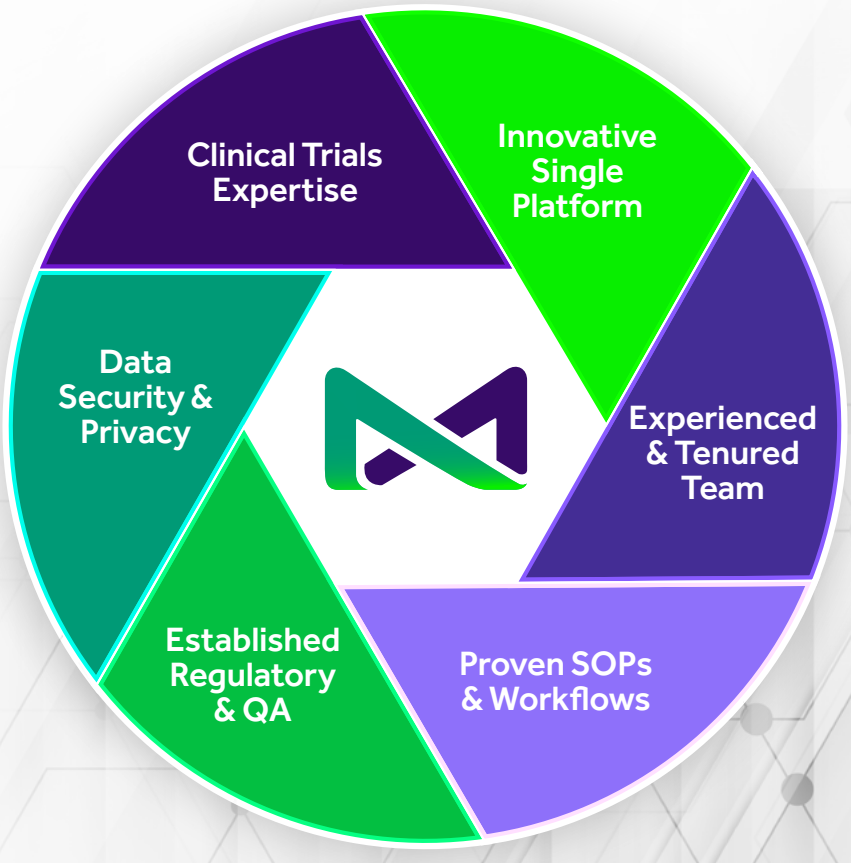
MERIT
Your Clinical Endpoint Expert

Oncology Imaging Solutions

MERIT has provided image collection, centralized reading, and data management services for clinical trials since 2012.

Through innovative and proven technologies, alongside our intuitive, seamless workflows, MERIT's experienced staff bring more than a decade of clinical endpoint expertise, ensuring the success and integrity of your clinical trials.

You have choices for your endpoint services provider. MERIT values your partnership and will make your study a priority.



ONCOLOGY TEAM EXPERIENCE:

- Team has managed 100+ oncology studies and across 20+ cancer types
- Team has cumulative experience of 40+ studies using RECIST 1.1 and other criteria, across all phases of clinical research
- Extensive experience among our highly qualified readers, KOLs, and medical advisor team

THE VALUE OF OUR PEOPLE

MERIT provides added value to complex clinical trial projects through personalized service, deep scientific expertise, and exceptional quality. Our team includes:

- Tenured project managers
- Dedicated imaging specialists
- Meticulous quality assurance staff
- Expert data managers and data quality evaluators
- Adept software developers

OUR READERS HAVE EXTENSIVE KNOWLEDGE AND TRAINING

- Our board-certified readers are dedicated academic radiologists, world leaders in their respective fields, and are able to read large volume of exams in a timely manner
- Participated in >1600 clinical trials and have contributed to >300 FDA or EMEA marketing authorization filings so far
- Trained in RECIST 1.1, iRECIST, mRECIST, Lugano, WHO, RANO, Hallek, PCWG3, Cheson, PERCIST, etc.
- Reading all modalities: MRI, CT, PET, SPECT & X-rays

EXCELSIOR™

EXCELSIOR is the **integrated, market-leading platform** designed for managing clinical trials that offers access to real-time data 24/7 for sponsors and clinical sites.

KEY EXCELSIOR FEATURES

- **Single System**
  One validated, secured system that contains all study data: images, tasks, queries, site materials, analysis, and read data
- **Universal**
  One cloud-based platform bypasses site IT challenges and effectively manages access rights for sites, sponsors, and readers (zero footprint implementation)
- **Transparent**
  Eliminates the question: *"where is our data?"* and reduces risk and data delays
- **Trusted**
  FDA 510k Medical Device

DOCUMENT PORTFOLIO MANAGEMENT

Study Design
- Expertise in reading criteria
- Imaging design/clinical data collection
- Clinical endpoints and surrogates

Quality Assurance Documentation
- Imaging Charter
- SOPs and robust learning management system
- Reader QA and monitoring

Standardized Imaging Procedures
- Provide imaging guidelines
- Monitor scan collection
- Query for issues

RECENT CASE STUDIES

Types of Savings

30%

Case Study I
- Phase I/IIa Retrospective Study of Glioma
- Response Criteria: RANO HGG/LGG

30% time reduction from resolving challenges with study start-up, read workflow, and data transfer

60%

Case Study II
- Phase II Newly Diagnosed GBM
- Response Criteria: RANO

60% reduction of cost associated with early phase Collect & Hold Project

(Internal data)

LEVERAGING OUR GLOBAL EXPERIENCE

TOTAL TRIALS HOSTED	>315
TOTAL SITES	>3,180
COUNTRIES	>59
TOTAL USERS	20,604
TOTAL PARTICIPANTS	57,298

COLLECT AND HOLD SERVICE

Our Collect and Hold service supports early-phase study image collection, quality control, cataloging, and storage in preparation for a potential read. This process can positively reduce start-up time for your projects. As your project progresses, MERIT can support additional capabilities including randomized reader and adjudication services.

MERIT

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Connect with us to learn more about how our expertise and approach can support bringing your product to market on-time and on-budget. Your success is our priority.

START A CONVERSATION

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