

MERIT: Bench-to-Bedside Comprehensive Endpoint Services, Tenured Medical Experts, EXCELSIOR™, AI-based Toolkit, Unmatched Global Footprint



Clinical Trial Endpoint Expertise

Dr. Ron Danis, MERIT's CSO, has 20+ years in leading clinical trial endpoint assessment and reading centers. MERIT's reading center currently has 30+ readers with the majority being ophthalmologists.



Comprehensive Software Tailored for Ophthalmic Trials

Best in class, user friendly, and most established cloud-based data system, EXCELSIOR™. First FDA 510(k) cleared platform provides the current gold standard in improved accuracy and efficiency.



Innovative Endpoint Assessment with Structural-functional Correlations

Quantitative endpoint assessment including machine learning tools for multiple retinal indications including GA region, EZ loss area, and other structural-functional correlations.



Operational Excellence

Processes and workflows are seamlessly controlled and managed through EXCELSIOR (MERIT's operational workhorse).



Transparency of Workflow

EXCELSIOR allows complete transparency of study data with 24/7 global access, all within one comprehensive system.



Extensive Global Footprint

MERIT has the largest world-wide experience, managing clinical sites in over 60 countries.



Preclinical IND-Enablement Offering

OSOD provides experienced leadership & operational capabilities for all facets of early stage development in ophthalmology.



Venture Partner Opportunity

Through its venture partner, Mianus Capital, MERIT can financially foster Sponsor innovation.

Comprehensive Ophthalmology Endpoint Services

MEDICAL & SCIENTIFIC EXPERTS

MORPHOLOGY

- OCT/OCTA
- FAF
- Color FP
- FA/ICGA
- Slit Lamp Imaging
- Specular Microscopy
- B Scan Ultrasonography
- IOL Rotational Analysis

R&D: NOVEL
ENDPOINTS

FUNCTION

- BCVA/LLVA
- Microperimetry
- Electroretinogram
- Visual Evoked Potential
- Pupillometry
- MLMT
- Light/Dark Adaptation
- Static & Kinetic Perimetry
- Contrast Sensitivity
- Ocular Motor Instability

EXCELSIOR™

FDA 510(k) Cleared Device

PATIENT DATA

Sponsors

Sites

CROs