

Specialized CRO for Hemato-Oncology Research

Robust Global Footprint: Presence Across 26 Geographies
spanning: **Americas | Asia-Pacific | Europe**



25+

Hemato-Oncology Trials



4500+

Patients Enrolled



550+

Sites

Supporting Hemato-Oncology Trials Through Our Multi-layered Approach

Treatment Approaches

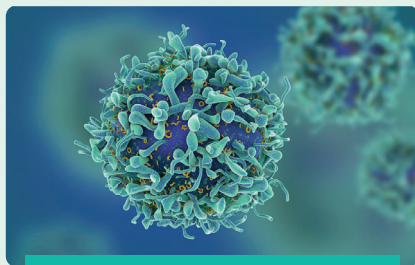
- Maintenance Therapy in Multiple Myeloma
- Innovative MRD Techniques
- Advancing Combination Therapy for RRMM
- Advancing Frontline Therapy for Transplant-Ineligible NDMM
- Innovative Therapies in Amyloidosis

Types of Treatments & Drug Classes

- Bispecific T-cell Engagers
- Antibody-Drug Conjugates
- Monoclonal Antibodies
- Proteasome Inhibitors
- Immunomodulatory Drugs

Hemato-Oncology Indications (Disease Focus)

Immune Thrombocytopenia | Waldenström's Macroglobulinemia | Hodgkin Lymphoma | AL Amyloidosis | Multiple Myeloma



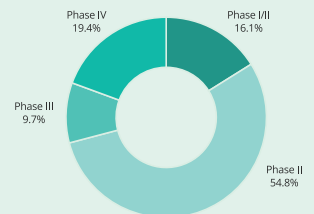
Global Leadership in Multiple Myeloma: Currently Supporting one of the largest trials

Full Service Capabilities

- Regulatory Affairs & Study Design
- Site Management & Patient Recruitment
- Clinical Monitoring & Data Management
- Pharmacovigilance & Safety Oversight
- Bioanalytical & Laboratory Services
- Project Management & Compliance

Scientific Capabilities

- Expertise in disease heterogeneity and MRD end points for tailored trial designs
- Adaptive trial designs ensuring efficiency without compromising data quality
- Specialized medical monitoring and PV teams for safety oversight and compliance
- Strong medical writing capabilities for regulatory and scientific documentation



Collaborated with the most renowned KOLs in designing and monitoring clinical trials

AI driven analytics platform to triage healthcare data for efficient trial design & execution

Real-time patient matching | EMR analysis for patient identification | Faster recruitment

The Veeda Lifesciences Advantage:

1

Faster Patient Recruitment

- Accelerate recruitment with our pre-vetted hemato-oncology patient database and strong site relationships
- Collaboration with centres and patient registries to identify eligible patients promptly

2

Optimized Budgets, Reliable Results

- Optimized site selection and adaptive trial designs to reduce costs
- Transparent pricing and tailored trial strategies to minimize cost overruns

3

Simplifying Regulatory Complexity

- Dedicated regulatory specialists & customized strategies to navigate globally and accelerate approval timelines

4

Robust Data Quality

- Statistics & analytics team ensuring continued data integrity

Integrated, Tech-Driven, Global CRO



Global Offices

Greece | India | Ireland | Italy



Scan for Case Studies



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Excellence
In Everything