



Veeda Oncology Network

An Experienced Network for Conducting Oncology Clinical trials

Pharmaceutical and Biotech companies face many challenges in conducting oncology clinical trials in today's research environment. The primary challenges are meeting stringent accrual timelines and obtaining partners that provide superior performance.

To enhance success of a clinical trial, sponsors should rely on an experienced, respected, and specialized clinical research partner, such as the Veeda Oncology Network with access to large pools of quality oncology research subjects to meet aggressive accrual goals. In addition, we are proven experts in managing clinical trial performance within our network sites which has been successfully demonstrated since our inception.

Distinguishing Features

Veeda Oncology functionality is supported by a group of professionals with extensive clinical trial knowledge. On average, each team member has an average of 15 years of Oncology specific experience. We provide the following services that benefit our pharmaceutical and biotech clients. These benefits have been demonstrated over the past 7 years, through more than 55 trials, equating to greater than 1,400 patients throughout our 350+ research qualified physicians and their qualified research staff.

Trial Design and Assessment

- Disease specific committee review
- Research Advisory Board review
- Investigator Initiated trial design
- Timely feasibility feedback

Trial Start-up

- Master Clinical Research Agreement (MCRA) on file for each site
- Site selection support
- Site specific profile information provided upon request
- One budget and contract for all participating sites
- Central regulatory management

Trial Operations

- Project management
- Site management
- Financial management
- Training
- Quality audits

Diseases Treated through Trial Experience

Veeda Oncology Network has developed a strong reputation in the industry as a result of our trial experience in the following disease types:

- Breast
 - Adjuvant ER+
 - 1st and 2nd Line Metastatic, ER/PR-, HER2Nu- /HER2Nu+
 - 1st/2nd Line Metastatic, ER/PR+, HER2Nu+
 - Supportive Care
 - Antiemetic
 - Bone Mets
- Chronic Lymphocytic Leukemia (CLL)
- Colorectal
 - 1st Line metastatic
 - 2nd Line metastatic
 - Supportive Care – antiemetic
- Hormone Refractory Prostate Cancer (HRPC)
 - 1st Line metastatic
 - 2nd line metastatic
 - Expanded access
- Lung - NSCLC
 - 1st Line metastatic
 - 2nd line metastatic
 - 3rd line metastatic
 - Non-adeno
 - Non-squamous
 - Anemia associated with NSCLC
- Supportive care (thrombocytopenia in NSCLC; antiemetic)
- Lung – SCLC
 - 1st line metastatic
 - 2nd line metastatic
- Mantle Cell Lymphoma (MCL)
- Myelodysplastic Syndrome (MDS)
 - 1st Line
- Melanoma
 - 1st Line
 - 2nd line
- Multiple Myeloma
- Non-Hodgkin's Lymphoma (NHL)
- Ovarian
 - 2nd Line
- Pancreatic
 - 1st Line
- Prostate
 - 1st line
 - 2nd Line
 - Expanded Access
- Renal Cell
 - 1st Line

Benefits of Utilizing the Veeda Oncology Network

Each trial is carefully assessed for its scientific merit as well as its functionality in a community-based setting. The assessment is completed through a disease specific committee of oncologists, and a Research Advisory Board (RAB) composed of both physicians and research coordinators.

By utilizing the network, your start-up timelines will be decreased by structured and timely site feasibility, negotiating one contract/one budget for all participating network sites, and regulatory document management allowing for rapid site approval. Our success in assisting you in managing your trial from start-up through close out, is in large part due to our long standing relationships with our investigators and research coordinators. This increases response time be it communication, accurate regulatory/safety maintenance, data submission and query resolution.

For your next oncology clinical trial, we hope you consider utilizing the Veeda Oncology Network so you too can become a part of our portfolio of success stories.



For additional inquiries or questions, please contact:

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