

PAZOPANIB CLINICAL TRIALS: OVERVIEW OF VEEDA'S CAPABILITIES



Pazopanib

Pazopanib is the generic name and is a targeted therapy that is classified as a Tyrosine Kinase inhibitor; Vascular Endothelial Growth Factor inhibitor. Pazopanib is used for the treatment of advanced renal cell carcinoma.



Up to 30% of patients have metastases at the time of the initial diagnosis

Key Differentiators

What makes Veeda stand apart when it comes to conducting Pazopanib studies?



Knowledge and experience of the proposed sites



Highly experienced team and effective project management



Close collaboration and planning with sites for effective and rapid recruitment



Advanced technological data monitoring solution



Proactive planning to anticipate risk and develop sound mitigation strategies



Comprehensive understanding and communication strategy involving multiple stakeholders



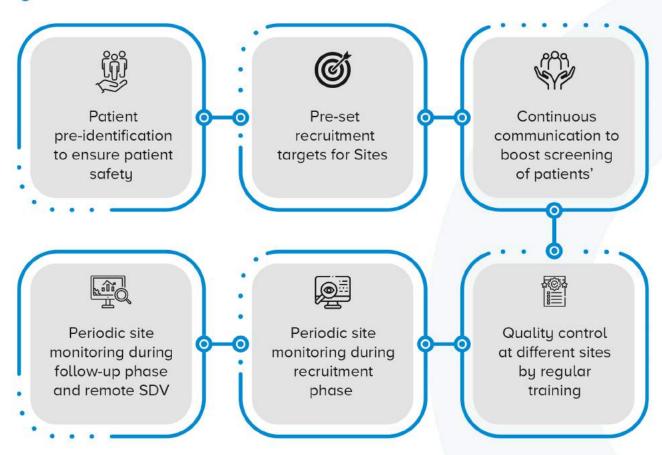
Quality is in our DNA

Proactive Quality and Compliance Control for Clinical Studies

- Aligned Sponsor and CRO Processes for the conduction of the study
- Detailed Quality Management Plan before starting with study for our quality client support
- Ensured for all internal pieces of training like GCP, Study-specific training, Mandatory protocol training with knowledge assessment to utilize our CRA Resource for quality study
- Study specific quality management plan and site visit by PM, as needed
- Mandatory Site Education: GCP training, Protocol training during IM, SIV, and Ongoing training at IMVs, Tools, and aids provided

Recruitment Plan for Success

Strategies for Better Patient Recruitment and Retention for a healthier tomorrow





Veeda Clinical Research

Your Capable, Knowledgeable and Reliable Partner for Pazopanib Trials

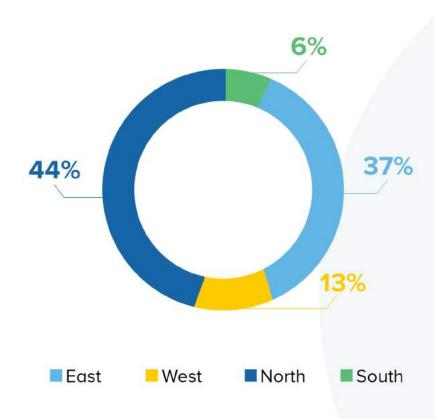
With a strong site network of 25+ experienced sites for Pazopanib study, our clinical operations team is accountable for project delivery, starting from developing projects plan to deciding timelines, to on-time delivery with ensured quality.



Our project management team carries out the following activities:

- Develop Project Plans, Timelines and Define Milestones
- Strategize with Functional Leads
- Oversee Site activities
- Communication with CRAs

Proposed Site Network in India





Streamline Your Clinical Trials

Gain complete control over your clinical trial operations with CTMS (Clinical Trials Management System):

- One centralized database for all study data
- Streamline monitoring visits
- Maintain electronic Trial Master File (eTMF)
- Automated email alerts/notifications
- Integrated IP Inventory tracking
- Real-time visibility into study milestones & metrics
- Track study budget & pass-through budget
- Automate investigator payments

Veeda's Expertise in Trials in Numbers!

- Database of 900+ Investigators
- Experience of working with 300+ Clinical Research Investigators
- Database of 150+ oncologists at 90 sites across India
- Conducted Patient PK, outpatient and hospitalized Oncology treatment studies in all stages of execution
- Data insights for Oncology recruitment :700+ Oncology patients enrolled

To know more about

our expertise in Clinical Trials, mail us at info@veedacr.com



www.veedacr.com