

Study to Learn More About the Safety and Effectiveness of the Drug VITRAKVI During Routine Use in Patients With TRK Fusion Cancer Which is Locally Advanced or Spread From the Place Where it Started to Other Places in the Body

NCT04142437

Status	RECRUITING
Sponsor	Bayer
Enrollment	150 participants

Key Eligibility Criteria

Inclusion (7)

- Adult and pediatric (from birth to 18-year-old) patients
 - Patients with locally advanced or metastatic solid tumor harboring an NTRK gene fusion. NTRK (NTRK1, NTRK2, and NTRK3) gene fusions will be identified locally. Acceptable methods of detection of NTRK gene fusion include NGS, fluorescence in situ hybridization (FISH), reverse-transcription polymerase chain reaction (rt-PCR) or any other genomic testing able to detect NTRK gene fusion. If a pan-TRK IHC method is used, this result needs to be accompanied with the results using one of the other methods noted above.
 - Life expectancy of at least 3 months based on clinical judgement
 - Decision to treat with larotrectinib made by the treating physician prior to study enrollment
 - Patients can also be enrolled if the initial visit (larotrectinib start date) occurred within 2 months \pm 3 days prior to informed consent signed date
- ... and 2 more (see full listing online)

Exclusion (5)

- Any contraindications as listed in the local approved product information
- Pregnancy
- Participation in an investigational program with interventions outside of routine clinical practice
- Prior treatment with larotrectinib or other kinase inhibitor with TRK inhibition
- Patients with NTRK gene amplification or NTRK point mutation

Locations (76 total)

Banner Desert Medical Center, Mesa, Arizona, United States
California Research Inst., Los Angeles, California, United States
USC / Norris Comprehensive Cancer Center, Los Angeles, California, United States
... and 73 more locations