Clinical Trial of VBC103 in Patients With Advanced Malignant Solid Tumors

NCT07299747

Status	RECRUITING
Phase	Phase 1, Phase 2
Sponsor	VelaVigo Bio Inc
Enrollment	255 participants

Key Eligibility Criteria

Inclusion (8)

- The subject or their legal representative is willing and able to sign a written ICF before initiating any study procedures.
- Histologically or cytologically confirmed unresectable advanced/metastatic solid tumor that has recurred or progressed during or after standard systemic therapy, or is intolerant to standard therapy, or lacks standard treatment options (applicable only to Phase I and Phase IIa Cohort 5).
- At least one measurable lesion as assessed by the investigator per RECIST v1.1.
- Adult male or female (defined as e18 years of age)
- ECOG performance status score of 0-1.

... and 3 more (see full listing online)

Exclusion (4)

- Any unresolved eGrade 2 toxicity from prior anticancer therapy.
- Known active keratitis or corneal ulcer.
- History of interstitial lung disease (e.g., non-infectious interstitial pneumonia, pneumonitis,pulmonary fibrosis, or severe radiation pneumonitis), current interstitial lung disease, or suspected interstitial lung disease based on imaging during the screening period.
- History of underlying pulmonary diseases, including but not limited to pulmonary embolism within 3 months prior to the start of investigational product, severe asthma, severe chronic obstructive pulmonary disease, restrictive lung disease, and other clinically significant pulmonary impairment or requiring supplemental oxygen, as well as any autoimmune, connective tissue, or inflammatory disease involving the lungs (such as rheumatoid arthritis, Sjögren's syndrome, sarcoidosis, etc.) and/or prior pneumonectomy (complete resection).

Locations (1 total)

Fudan University Shanghai Cancer Center, Shanghai, Shanghai Municipality, China