ForPatients
by Roche

A Study of TACE Combined With Atezolizumab Plus Bevacizumab or TACE Alone in Patients With Untreated Heaptocellular Carcionma

Trial Status	Trial Runs In	Trial Identifier
Active, not recruiting	2 Countries	NCT04712643 ML42612

The information is taken directly from public registry websites such as ClinicalTrials.gov, EuClinicalTrials.eu, ISRCTN.com, etc., and has not been edited.

Official Title:

A Phase III, Open-Label, Randomized Study of On-Demand TACE Combined With Atezolizumab Plus Bevacizumab (Atezo/Bev) or On-Demand TACE Alone in Patients With Untreated Heaptocellular Carcionma

Trial Summary:

This study will evaluate the efficacy and safety of atezolizumab plus bevacizumab combined with on-demand TACE compared to on-demand TACE alone in participants with hepatocellular carcinoma who are at high risk of poorer outcome following TACE treatment.

Hoffmann-La Roche
Sponsor

Phase 3
Phase

NCT04712643 ML42612
Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	#18 Years	No

Inclusion Criteria:

- Confirmed diagnosis of HCC by histology/ cytology or clinical criteria
- Eligible for TACE treatment
- No prior systemic therapy for HCC, especially immunotherapy
- No prior locoregional therapy to the target lesion(s)
- At least one measurable untreated lesion
- ECOG Performance Status of 0-1
- Child-Pugh class A

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Exclusion Criteria:

- Evidence of Vp3/4 and hepatic vein tumor thrombus (HVTT)
- Evidence of extrahepatic spread (EHS)
- Being a candidate for curative treatments
- Any condition representing a contraindication to TACE as determined by the investigators
- Active or history of autoimmune disease or immune deficiency
- Untreated or incompletely treated esophageal and/or gastric varices with bleeding or high risk for bleeding
- A prior bleeding event due to esophageal and/or gastric varices within 6 months prior to initiation of study treatment
- Evidence of bleeding diathesis or significant coagulopathy

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