

Clinical Trial to Evaluate the Efficacy and Safety of Cellgram-LC Administration in Patients With Alcoholic Cirrhosis

NCT04689152

Status	RECRUITING
Phase	Phase 3
Sponsor	Pharmicell Co., Ltd.
Enrollment	200 participants

Key Eligibility Criteria

Inclusion (7)

- At the time of screening, 19 or 70 years
 - Patients diagnosed with alcoholic cirrhosis by combining alcohol history, imaging and pathological examination results, and clinical symptoms at screening, and belonging to Child-Pugh grade B or C (Child-Pugh score of 7 or more)
 - Those whose survival period is more than 1 year when judged by the tester
 - Those who can perform hepatic artery catheterization by inserting a catheter into the hepatic artery at the judgment of the examiner
 - In the case of women of childbearing potential, a person who was confirmed negative in the pregnancy test at screening and agreed to use contraception* by the method permitted for this clinical trial during the clinical trial
- ... and 2 more (see full listing online)

Exclusion (25)

- Those with a history of solid cancer including Hepatocellular Carcinoma (HCC) (within 5 years before screening), those who have been diagnosed with solid cancer and are currently undergoing chemotherapy or those whose hepatocellular carcinoma has been confirmed by screening tests
 - Patients who underwent portal systemic shunting in the jugular vein
 - Patients with alcohol consumption or hepatotoxic drugs within 6 months prior to screening
 - Persons taking high-dose steroids, immunosuppressants, or antimicrobials due to severe infections for at least 1 month of screening
 - Those who have major surgical operations, long-term biopsy, or significant trauma as judged by the investigator within 3 months before screening
- ... and 20 more (see full listing online)

Locations (11 total)

Soonchunhyang University Hospital, Bucheon-si, South Korea
Soonchunhyang University Hospital, Cheonan, South Korea
Gangwon National University Hospital, Chuncheon, South Korea
... and 8 more locations

<https://clinicaltrials.gov/study/NCT04689152>

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