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Celltrion's solid tumor ADC drug candidate 'CT-P73' has received approval for phase 1 clinical trials in Korea.

🕆 Reporter Ji Yong-jun 📗 🗡 Posted 2025.11.10 15:42 📗 🗡 Last updated on November 10, 2025, at 3:45 PM . □Comments 0

Phase 1 clinical trial of CT-P73 monotherapy in adult patients with advanced solid tumors





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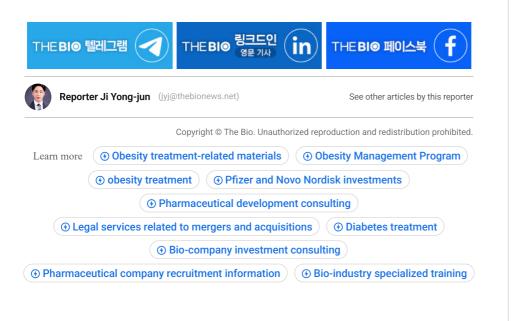
According to industry sources on the 10th, Celltrion received approval from the Ministry of Food and Drug Safety for its domestic Phase 1 clinical trial (Investigational New Drug) for CT-P73. This clinical trial will evaluate the safety, tolerability, pharmacokinetics, immunogenicity, and initial efficacy of CT-P73 monotherapy in adult patients with advanced solid tumors. Celltrion anticipates that the drug will be effective in cancers such as cervical, head and neck, and colorectal cancers.

CT-P73 is an ADC therapeutic candidate that targets a specific tissue factor highly expressed in cancer cells and induces a payload (cytotoxic anticancer drug) linked to an antibody via a linker to attack cancer cells. The company is developing CT-P73 as a "best-in-class" drug.

Through open innovation, Celltrion applied PBX-7016, a novel camptothecin-derived topoisomerase (TOP1) inhibitor payload jointly developed with domestic biotech company PinoBio, to CT-P73. TOP1 inhibitors are drugs that kill cancer cells by damaging DNA and inhibiting their spread.

Celltrion unveiled preclinical results for CT-P73 at World ADC in June. The company explained that CT-P73 achieved significant results in terms of the therapeutic index (TI), a comprehensive assessment of efficacy and safety, based on low toxicity and superior efficacy.

Furthermore, based on the proven stability of the linker, superior pharmacodynamic properties were confirmed. Furthermore, Celltrion expects that CT-P73 administration will demonstrate excellent tumor penetration as well as a bystander effect, destroying surrounding cancer cells.



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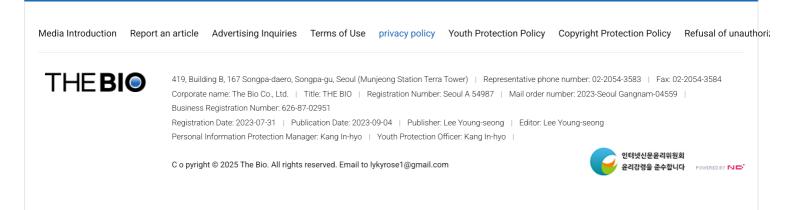
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