

## **Bukwang Pharma receives FDA approval for dyskinesia drug trial**

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**The approved trial will be conducted in about 30 institutions in the United States**



South Korea based Bukwang Pharmaceuticals recently announced that the US Food and Drug Administration (FDA) has approved a phase 2 trial of JM-010.

JM-010 is a new drug candidate developed by Danish Bioventure Contera Pharma, a subsidiary of Bukwang Pharmaceutical.

The approved trial will be conducted in about 30 institutions in the United States to evaluate the safety and effectiveness of JM-010 in 190 patients with side effects of dyskinesia caused by Parkinson's disease. Dyskinesia is uncontrolled, involuntary movement that may occur with long-term levodopa use and longer time with Parkinson's.

As JM-010 is entering the global clinical stage, it is already well-proven and plans to enter the KOSDAQ through the technology listing of foreign companies.