Completion of CMC, GLP-tox Testing, Clinical Trial Medication Production, and IND Approval for Orally Administerable CD39 Target Inhibitor Preclinical Candidate Substance



ONCOLOGY	Preclinical
Product Type	Small Molecule
Indication	Solid cancer, Pancreatic Cancer
Target	CD39
MoA(Mechanism of Action)	Potent anti-tumor effect through boosting of anti-tumoral immunity in Tumor-associated microenvironment
Competitiveness	Down stream target of ATP/ADO signal axis, CD73 inhibitors such as Quemlistat (clinical phase 1b) Superiority of CD39 as drug target compared to CD73
Development Stage	Currently under CMC development, In this year (2024), it will be entered to GLP-tox for 2025 clinical trial
Route of Administration	Oral administration

