

# Highlights from the Week of 30 June, 2025 - 07 July, 2025



Unicycive Therapeutics receives a complete response letter for oxylanthanum carbonate to treat hyperphosphatemia in dialysis patients with chronic kidney disease

*(FDA, CR)*



Takeda receives US FDA approval for GAMMAGARD LIQUID ERC, a ready-to-use low IgA therapy for primary immunodeficiency; the launch is planned for 2026, with the first-generation product to be discontinued

*(FDA Approval)*



Positive results from Amgen's Phase 3 trial showed that bemarituzumab plus chemotherapy significantly improves overall survival over chemotherapy alone in FGFR2b-positive gastric cancer

*(Clinical trial, Results)*

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FDA grants accelerated approval to Dizal's ZEGFROVY, marking it as the exclusive oral targeted treatment for EGFR exon 20 insertion non-small cell lung cancer

*(FDA Approval)*



FDA approves Hikma's TYZAVANTM, a ready-to-infuse vancomycin for conditions like skin and bone infections and lower respiratory tract infections

*(FDA Approval)*



European Commission approves Biocon Biologics' VEVZUO for preventing bone complications in advanced cancer and EVFRAXY for treating osteoporosis

*(Biosimilars, EC Approval)*

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Neurizon enters into a global license agreement with Elanco to accelerate the commercialization of NUZ-001, an investigational therapy for neurodegenerative diseases like ALS

*(Commercialization, Partnership)*



Jazz Pharmaceuticals' ZIIHERA (zanidatamab) receives European Commission approval as the EU's first HER2-targeted therapy for advanced HER2-positive biliary tract cancer

*(EU Approval)*