

Highlights from the Week of 1st January 2026 to 12th January 2026



FDA approves ScinoPharm's
injectable GLATIRAMER
ACETATE for multiple sclerosis

(FDA, Approval)



Biocon Pharma receives FDA
approval for EVEROLIMUS
tablets for oral suspension,
indicated for the treatment of
certain types of tumors and
tuberous sclerosis complex

(FDA, Approval)



European Commission
approves GSK's SHINGRIX
recombinant zoster vaccine in
a prefilled syringe
presentation

(EU, Approval)

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Complement Therapeutics gains Fast Track designation for its gene therapy CTX001, to treat geographic atrophy secondary to age-related macular degeneration

(FDA, Fast Track)



Oculis gains FDA Breakthrough Therapy designation for Privosegtor to treat patients with optic neuritis

(FDA, Breakthrough Therapy)



FDA grants Orphan Drug designation to Arbele's ARBI002, an anti-CDH17 antibody drug conjugate for pancreatic cancer treatment

(FDA, Orphan Drug)

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The Sanofi logo, consisting of the word 'sanofi' in a lowercase, bold, sans-serif font with a small purple dot above the 'i'.

Sanofi's sBLA for Tzielid in children aged one to seven with stage 2 type 1 diabetes receives FDA priority review

(FDA, Priority Review)

The Vera Therapeutics logo, featuring the word 'vera' in a lowercase, bold, sans-serif font with a blue-to-white gradient, and 'therapeutics™' in a smaller, lowercase, sans-serif font below it.

Vera Therapeutics gains priority review for its BLA atacicept in IgA nephropathy, with a PDUFA target action date of June 3, 2026

(FDA, Priority Review)

The Novo Nordisk logo, featuring a stylized cow icon above the text 'novo nordisk®' in a lowercase, sans-serif font.

Novo Nordisk makes the WEGOVY pill available in the US as the first and only oral GLP-1 for adult weight management and cardiovascular risk reduction

(Commercialisation)