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**Oncolys BioPharma Inc.**

# Announcement of Completion of Enrollment in Phase IIa Clinical Trial of OBP-601 (censavudine, TPN-101) in Patients with Progressive Supranuclear Palsy (PSP)

Oncolys BioPharma Inc. (hereafter "Oncolys") today announced that it has received a notice from Transposon Therapeutics, Inc. ("Transposon"), licensee of OBP-601 (censavudine, TPN-101), that enrollment has been completed with 42 patients randomized in the Phase IIa, double-blind, placebo-controlled study of OBP-601 in patients with Progressive Supranuclear Palsy (PSP). The target date for publication of the clinical trial results has not yet been determined.

In addition, the Phase IIa, double-blind, placebo-controlled study in patients with Amyotrophic Lateral Sclerosis (ALS) and/or Frontotemporal Degeneration (FTD) is recruiting patients.

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**About Oncolys BioPharma Inc.**

Oncolys BioPharma Inc.(JPX:4588) develops novel cancer therapeutics and diagnostic products using gene modified viral technologies and aims to contribute to fulfill unmet medical needs for cancer and severe infectious diseases. Especially in oncology area, we utilize technology platform for oncolytic virus and develop Telomelysin and its next-generations for cancer treatment and TelomeScan for early detection of cancer and recurrence monitoring after surgery. We have established broad range of product pipeline to cover early detection of cancer, early treatment of local cancer, post-operative examination, and treatment of metastatic cancer. For more information, please visit http://www.oncolys.com/en/

Contact:
**Oncolys BioPharma Inc.**
Email: oncolys_information@oncolys.com