

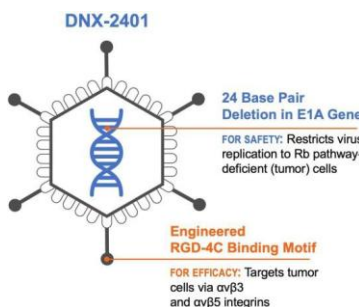


DNX-2401

Investment Opportunity

Differentiated Intratumoral Oncolytic Adenovirus with Dual Oncolytic + Immune-Stimulatory Activity

- Engineered adenovirus: 24 bp E1A deletion (tumor-selective replication) + RGD-4C motif for integrin targeting
- Intratumoral delivery limits systemic exposure
- Strong clinical signals:
 - Pediatric Ph. 1 monotherapy in DIPG: median OS = 17.8 months (durable responses vs historical SOC); ORR 25% associated with longer survival
 - Adult Ph. 1 monotherapy in recurrent glioblastoma multiforme: median OS = 9.5 months
- Safety profile: low systemic toxicity and manageable intratumoral administration in pediatrics and adults
- Regulatory tailwinds: FDA Fast Track, orphan and rare pediatric disease designations; priority review voucher eligibility for DIPG pathway
- Uniquely positioned as potential first-line monotherapy in pediatric DMG



Clinical Partners

FDA DIPG Rare Pediatric Designation
FDA DIPG Fast Track
FDA Orphan Drug & Fast Track Designation
EU Orphan Drug Designation

Proposed Convertible Note Offering

CV Bio is raising an \$8.0M Convertible Bridge Note to fund operations through a potential strategic transaction within 12–18 months

CV Bio presents a compelling investment opportunity in DNX-2401, a clinically validated, intratumoral oncolytic adenovirus immunotherapy uniquely positioned to address high-unmet-need pediatric brain cancers — including Diffuse Intrinsic Pontine Glioma (DIPG), Diffuse Midline Glioma (DMG), and recurrent pediatric high-grade gliomas. To advance this program, CV Bio is raising an \$8,000,000 Convertible Bridge Note to fund operations through a potential strategic transaction within the next 12 to 18 months.

Attractive profile driving strategic interest:

- Two active international pediatric brain tumor trials underway
- Clear DIPG pathway toward accelerated approval
- FDA Priority Review Voucher eligible
- Robust Ph. 1 clinical data in DIPG and currently advancing across international Ph. 1/2 and Ph. 2 studies evaluating optimized repeat-dosing regimens and expanded pediatric indications

Upcoming Milestones

- 2026**
 - Q1'26: Initiate Ph. 1/2 PNOc DMG Trial
 - Q2'26: New tumor types initial safety data; Dose escalation safety data
 - Q3'26: Repeat dose feasibility data
 - Q4'26: Initiate expansion phase
- 2027**
 - 1H'27: FDA meeting to discuss accelerated approval with expansion

Convertible Bridge Note Terms

- Up to \$8.0M aggregate principal amount
- Accrues interest at 8.0% per annum, with a 20% discount (mandatory conversion at 80%) of the purchase price in an M&A transaction
- Matures in 3 years. If not converted sooner, it converts at maturity into Series A shares at \$1.00 per share

Please direct all inquiries regarding this transaction to:

Harry Bushong
President

hbushong@convergence-ventures.com