

Palisades Therapeutics Unveils Breakthrough Multi-Mechanistic Data for PT150 and PT157 in Resistant Cancers

Dual AR/GR Antagonism, Metabolic Reprogramming, and Lysosomal Bypass Position PT150 and PT157 as Best-in-Class Oncology Candidates

CLIFFSIDE PARK, NJ, UNITED STATES, April 24, 2025 /EINPresswire.com/ -- Pop Test Oncology LLC dba [Palisades Therapeutics](#), a clinical-stage biopharmaceutical company, today released compelling preclinical data for clinical-stage PT150 and its early-stage dimer PT157, demonstrating their superiority over existing glucocorticoid receptor (GR) antagonists like Relacorilant™.

These findings position PT150 and PT157 as transformative therapies for taxane-resistant and AR/GR-driven cancers, including pancreatic, ovarian, liver, prostate, and colorectal malignancies.

Breakthrough Mechanistic Insights

- **Dual AR/GR Targeting:** Disrupts survival pathways in AR+/GR+ tumors (~30% of pancreatic, ovarian, and prostate cancers).
- **Epigenetic Modulation:** Reverses PARP resistance (H2AFV \square) induces synthetic lethality in ARID1A-mutant tumors (SMARCA4 \square).



Pop Test/Palisades Therapeutics

Key Differentiators: PT150/PT157 vs. Relacorilant™ according to Perplexity ai

Parameter	PT150/PT157	Relacorilant™
Mechanism	Dual AR/GR blockade + epigenetic/metabolic modulation + lysosomal bypass	GR antagonism only
Target Pathways	- Epigenetic (H2AFV \uparrow , SMARCA4 \downarrow) - Metabolic (ACADL \downarrow , SDHA \uparrow) - Stromal (ACTA2 \downarrow) - Lysosomal (RAB9A \downarrow)	GR-SGK1/DUSP1 pathway only
Preclinical Efficacy	- 55.3% TGI in taxane-resistant breast/ovarian cancer - 100% survival in PDX ovarian models - IC50 advantage in resistant pancreatic/liver cancer cells - ZTX® PREDICT in Vivo Platform Validation: Synergy with standard care in prostate models	60.5% TGI (pancreatic models)
Combination Potential	Synergizes with taxanes (Abraxane/Cabazitaxel), PARP inhibitors, HSP90 modulators	Limited to nab-paclitaxel
Biomarkers	RAB9A \downarrow , SMARCA4 \downarrow , KALRN \downarrow , OVOL2 \uparrow	SGK1 \downarrow , DUSP1 \downarrow
Market Potential	Targets AR+/GR+/ARID1A-mutant tumors across \$8B+ indications	\$2B niche (GR-high ovarian cancer)

Key Differentiators: PT150 and PT157 vs. Relacorilant

- Metabolic & Stromal Targeting: ACADL \square disrupts fatty acid oxidation; ACTA2 \square reduces fibrosis for improved drug delivery.

- Lysosomal Resistance Bypass: RAB9A \square prevents taxane sequestration, enhancing chemotherapy efficacy.

- ZTX[®] PREDICT Platform Validation: Rapid 5-day in vivo testing identifies responders/non-responders, with high prediction of patient treatment outcome, accelerating preclinical-to-clinical translation.

Strategic Next Steps

- Global Partnerships: Leverage data for potential collaborations with leading companies such as Pfizer, Gilead, Bristol-Myers Squibb, and Astellas.

Compelling Preclinical Data Across Multiple Cancers

Cancer Type (Site)	Model	Treatment	Key Findings
Pancreatic Cancer (Georgetown University)	n-PTX-resistant PDAC	PT150/PT157 + nab-paclitaxel	Overcomes resistance (IC50 advantage); RAB9A \downarrow , ACTA2 \downarrow , SMARCA4 \downarrow , H2AFV \uparrow
Ovarian Cancer (TD2/TGEN, Sai Life Science)	PDX models	PT157 + Enzalutamide	100% survival at 60 days; robust synergy in AR+/GR+ subtypes
Triple-Negative Breast Cancer (TD2/TGEN)	Xenograft	PT150 + Paclitaxel	55.3% tumor growth inhibition (TGI); 2.7x tumor volume reduction
Liver Cancer (HCC/CHC) (Kurume University-Japan)	11 HCC & 2 CHC cell lines	PT150/PT157	Cell growth suppressed in 10/11 (PT150) and 11/11 (PT157) lines; IC50 5.1–19.8 μ M
Prostate Cancer (Mount Sinai & BioReperia)	Zebrafish Tumor Xenograft	PT150/PT157 + Standard Care, Enzalutamide	Synergistic inhibition of metastasis; rapid 5-day efficacy testing
Prostate Cancer (Mount Sinai & Sai Life Science)	CRPC cell lines	PT150/PT157 + Abiraterone/ARN509/Enzalutamide	6.8x boost in cell death (PT150); PT157 2–4x more potent than AR antagonists

Compelling Preclinical Data Across Multiple Cancers

Why This Matters: PT150 and PT157’s multi-target approach offers hope for durable remissions in resistant pancreatic, ovarian, liver, and prostate cancers.

[About Palisades Therapeutics](#)

Palisades Therapeutics is a lean, innovation-driven biotech leveraging a global network of 75+ experts to accelerate transformative oncology therapies. Learn more at www.palisadestherapeutics.com

Disclaimer:

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Forward-looking statements: This release contains projections based on preclinical data. Clinical outcomes may differ.

#Pfizer, #Gilead, #BMS, #Astellas, #Oncology, #CancerResearch

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