



SUVN-D4010

5-HT₄ Partial Agonist

Dual Mechanism of Action (Disease Modifying and Symptomatic Treatment Potential for AD)

Phase 1 Completed in USA; Ready for Phase 2 POC

SUVN-D4010: Key Features

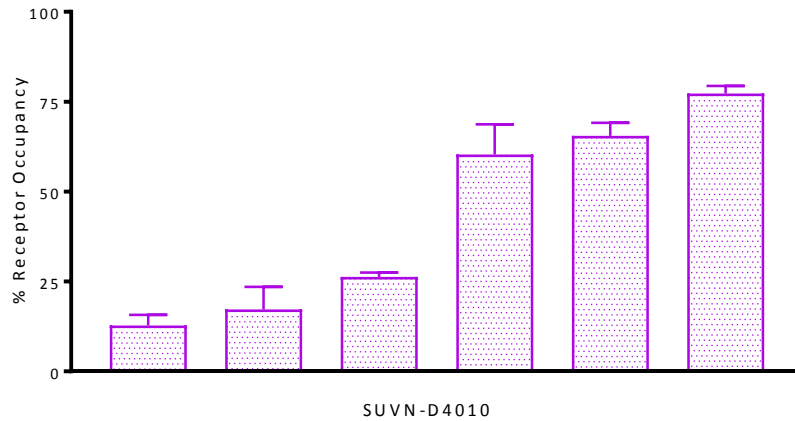


- Selective serotonin-4 receptor (5-HT₄) partial agonist
- Orally bioavailable with good brain penetration
- Good correlation between affinity, free fraction and efficacy
- Clean in hERG, phospholipidosis and AMES assays
- No QT/QTc prolongation in dog telemetry study
- Robust efficacy in animal models
- Potentiates the efficacy of cholinesterase inhibitors in animal models
- Disease modifying effects in animal models
- Translatable biomarker available for POC study
- Good margin of safety in preclinical studies
- No genotoxicity and teratogenic potential
- Well protected intellectual property in all major markets
- Safe and well tolerated in healthy adult subjects (Phase-1 study)
- Excellent human pharmacokinetics suitable for once a day treatment
- Projected human efficacy concentrations achieved in Phase-1 study
- No effect of food, gender and age on human pharmacokinetics

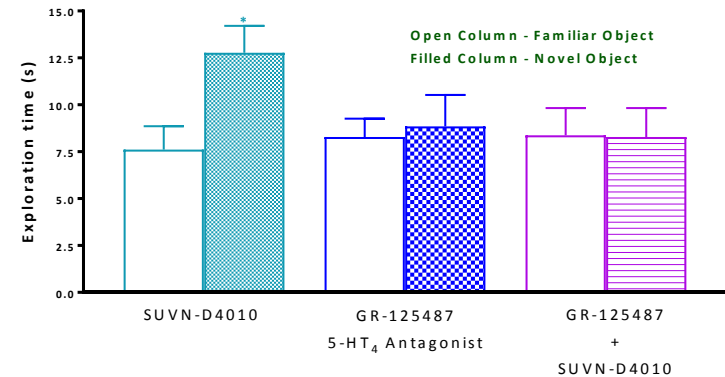
SUVN-D4010: Key Pharmacology Results



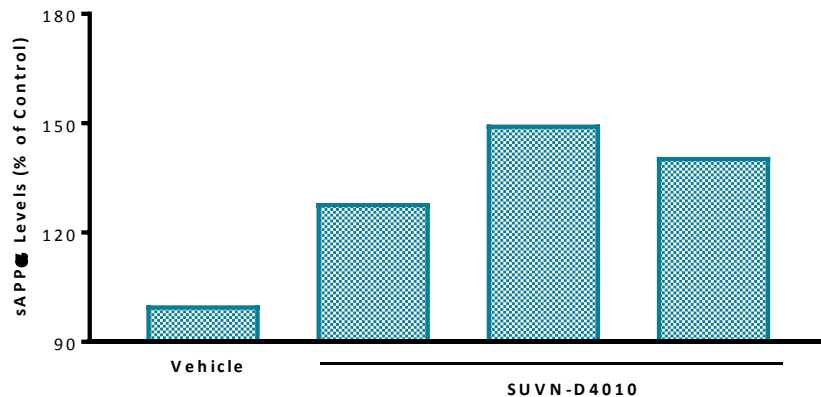
Target Engagement in Rats



Object Recognition Task in Rats



Modulation of sAPP α Levels



Disease Modifying and Symptomatic Treatment Potential for Alzheimer's Disease

SUVN-D4010: Clinical Studies Summary and Current Status



- Well tolerated in humans with dose dependent pharmacokinetics
- Suitable for once a day oral dosing
- Projected human efficacy concentrations achieved at low doses in Phase 1 study

Current status: Phase 1 Completed in USA; Ready for Phase 2 POC



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